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Access DB11 15-03.40

Scientific and Technical Information Center

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3 (US20040096496 OR US20040033262)/PN L2 L3

3 IN2002-MA601#/AP, PRN

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FILE COVERS 1907 - 15 Apr 2005 VOL 142 ISS 17 FILE LAST UPDATED: 14 Apr 2005 (20050414/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN

2004:412577 HCAPLUS AN

140:395542 DN

Entered STN: 21 May 2004 ED

Sustained release pharmaceutical composition of a cephalosporin antibiotic ΤI

Kshirsagar, Rajesh Suresh; Boldhane, Sanjay Parbhatrao; Jindal, Kour Chand Orchid Chemicals & Pharmaceuticals Limited, India IN

PA

S<sub>0</sub> U.S. Pat. Appl. Publ., 21 pp., Cont.-in-part of U.S. Ser. No. 222,930. CODEN: USXXCO

DT Patent

English LA

ICM A61K031-545 IC

ICS A61K009-20; A61K031-736

424465000; 514200000; 514054000 NCL

63-6 (Pharmaceuticals)

FAN.	CNT 3 PATENT NO.	KIND	DATE	APPLICATION NO.	DATE			
PΙ	US 2004096496	A1	20040520	US 2003-642194	20030818 <			
	US 2004033262	A1	20040219	US 2002-222930	20020819 <			
	WO 2004016250	A1	20040226	WO 2002-IB3320	20020819			
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PRAI IN 2002-MA601
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                                  20020819
     US 2002-222930
                           A2
     WO 2002-IB3320
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CLASS
 PATENT NO.
                  CLASS
                         PATENT FAMILY CLASSIFICATION CODES
 US 2004096496
                  ICM
                         A61K031-545
                         A61K009-20; A61K031-736
                  ICS
                         424465000; 514200000; 514054000
                 NCL
                         A61K031/545; A61K031/546
 US 2004096496
                  ECLA
 US 2004033262
                         A61K009/20H6B; A61K009/20H6F; A61K031/545; A61K031/546;
                  ECLA
                         A61K031/736
     Sustained release pharmaceutical compns. comprising at least a
     cephalosporin antibiotic, a mixture of polymers comprising of galactomannans
     and neutral swellable polymers, and other pharmaceutically acceptable
     excipients are described. The composition comprises about 30% to about 90% by
     weight of a cephalosporin antibiotic; about 1% to about 30% by weight of said mixture of polymers comprising from about 0.1% to about 15% by weight of
     galactomannans, and about 0.1% to about 15% of neutral swellable polymer
     by weight of sustained release composition
     cephalosproin antibiotic sustained release tablet
IT
     Drug delivery systems
        (carriers; sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
IT
     Lubricants
        (pharmaceutical; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
     Swelling agents
        (polymers; sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
IT
     Antibiotics
     Binders
     Drying
     Mixing
     Plasticizers
     Sieving
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     Gelatins, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
IT
     Polyesters, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
IT
     Drug delivery systems
        (sustained-release; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
     Polymers, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (swellable; sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
     Drug delivery systems
        (tablets, coated; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
     Drug delivery systems
        (tablets; sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
IT
     Fats and Glyceridic oils, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
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(Biological study); USES (Uses)
         (vegetable, hydrogenated; sustained-release pharmaceutical composition of a
         cephalosporin antibiotic)
     7631-86-9, Silicon dioxide, biological studies
ΙT
      RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
         (colloidal; sustained-release pharmaceutical composition of a cephalosporin
         antibiotic)
      9004-34-6, Cellulose, biological studies
      RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
      study); USES (Uses)
         (microcryst.; sustained-release pharmaceutical composition of a
         cephalosporin antibiotic)
     9005-25-8, Starch, biological studies
RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
      study); USES (Uses)
         (pregelatinized; sustained-release pharmaceutical composition of a
         cephalosporin antibiotic)
     9003-39-8, Polyvinylpyrrolidone 9004-62-0, Hydroxyethylcellulose
      9004-64-2, Hydroxypropyl cellulose
                                              9004-65-3, Hydroxypropyl
      methylcellulose
      RL: MOA (Modifier or additive use); POF (Polymer in formulation); THU
      (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     57-11-4, Stearic acid, biological studies 57-50-1, Sucrose, biological
IT
     studies 557-04-0, Magnesium stearate 557-05-1, Zinc stearate 1592-23-0, Calcium stearate 9000-30-0, Guar gum 11138-66-2, Xanthan
     gum 14807-96-6, Talc, biological studies
RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
IT
      153-61-7, Cephalothin 11078-30-1, Galactomannan
                                                               11111-12-9D,
                                   15686-71-2, Cephalexin
35607-66-0, Cefoxitin
                                                              21593-23-7, Cephapirin
50370-12-2, Cefadroxil
      Cephalosporins, derivs.
      34444-01-4, Cefamandole
     53994-73-3, Cefaclor 61270-58-4, Cefonicid 64544-07-6, Cefuroxime axetil 68401-81-0, Ceftizoxime 87239-81-4, Cefpodoxime proxetil
      92665-29-7, Cefprozil 117467-28-4, Cefditoren pivoxil
      RL: PEP (Physical, engineering or chemical process); PYP (Physical
      process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
      USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     9004-32-4
                   9010-88-2, Ethyl acrylate-methyl methacrylate copolymer
     9050-04-8, Calcium carboxymethyl cellulose 25086-89-9, Plasdone S-630 RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     50-70-4, Sorbitol, biological studies 50-99-7, Glucose, biological studies 63-42-3, Lactose 69-65-8, Mannitol 69-79-4, Maltose
      9004-53-9, Dextrin
                            66828-18-0, Dextrate
                                                      74811-65-7, Croscarmellose
      sodium
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN
      2004:162583 HCAPLUS
AN
      140:205146
DN
     Entered STN: 29 Feb 2004
      Sustained release pharmaceutical compositions of a cephalosporin
TI
      Kshirsagar, Rajesh Suresh; Boldhane, Sanjay Parbhatrao; Jindal, Kour Chand
IN
     Orchid Health Care, A Division of Orchid Chemicals & Pharmaceuticals Ltd.,
PA
      India
SO
     PCT Int. Appl., 55 pp.
      CODEN: PIXXD2
DT
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     English
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     ICM A61K009-22
IC
           A61K031-545; A61K031-546
      63-6 (Pharmaceuticals)
      Section cross-reference(s): 1
FAN. CNT 3
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PATENT NO.
                            KIND
                                    DATE
                                                 APPLICATION NO.
                                                                           DATE
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CLASS
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                           A61K031-545; A61K031-546
                           A61K009/20H6B; A61K009/20H6F; A61K031/545; A61K031/546;
 US 2004033262
                   ECLA
                           A61K031/736
AB
     Sustained release pharmaceutical compns. comprising at least a
     cephalosporin, a mixture of polymers comprising of galactomannans and
     neutral swellable polymers, and other pharmaceutically acceptable excipients are described. The composition comprises 30-90% of a cephalosporin
     antibiotic, 1-30% mixture of polymers comprising 0.1-15% galactomannans, and
     0.1-15% by weight of a neutral swellable polymer. Thus, tablets contained cephalexin monohydrate 75.73, lactose 6.27, xanthan gum 7.0, Eudragit NE
     30D 7.0, HPMC E5 31.5, Mg stearate 1.0 and water qs to 100%.
     sustained release cephalosporin polymer
     Drug delivery systems
         (granules, sustained release; sustained release pharmaceutical compns.
         of cephalosporins)
IT
     Binders
     Dissolution
     Drug bioavailability
     Human
     Lubricants
     Plasticizers
         (sustained release pharmaceutical compns. of cephalosporins)
     Carbohydrates, biological studies
IT
     Gelatins, biological studies
     Polymers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained release pharmaceutical compns. of cephalosporins)
IT
     Drug delivery systems
         (sustained-release; sustained release pharmaceutical compns. of
         cephalosporins)
IT
     Drug delivery systems
         (tablets, sustained-release; sustained release pharmaceutical compns.
         of cephalosporins)
     Fats and Glyceridic oils, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
ΙT
         (vegetable, hydrogenated; sustained release pharmaceutical compns. of
         cephalosporins)
     9004-34-6, Cellulose, biological studies
IT
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RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (microcryst.; sustained release pharmaceutical compns. of
         cephalosporins)
     15686-71-2, Cephalexin 50370-12-2, Cefadroxil
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     study); USES (Uses)
         (sustained release pharmaceutical compns. of cephalosporins)
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                           9000-30-0, Guar gum 9000-40-2, Locust bean gum
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Hydroxyethyl cellulose 9004-64-2, Hydroxypropyl cellulose
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     9010-88-2, Eudragit NE 30D 9050-04-8, Calcium carboxymethyl cellulose
     9063-38-1, Sodium starch glycolate 11078-30-1, Galactomannan
     11111-12-9, Cephalosporin 11138-66-2, Xanthan gum 14807-96-6, Talc, biological studies 21593-23-7, Cephapirin 23325-78-2, Cephalexin
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         (sustained release pharmaceutical compns. of cephalosporins)
RE. CNT
               THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
(1) Lupin Lab Ltd; WO 0241876 A 2002 HCAPLUS
    Shankar, M; WO 02067943 A 2002 HCAPLUS
(2)
(3) Squibb Bristol Myers Co; WO 9846213 A 1998 HCAPLUS
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     Sustained-release pharmaceutical composition of a cephalosporin antibiotic
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                   ECLA
                          A61K031/545; A61K031/546
 US 2004096496
     A sustained release pharmaceutical composition comprising at least a
     cephalosporin antibiotic and a mixture of polymers and other
     pharmaceutically acceptable excipients is described. Polymers are
     selected from mixture of galactomannans and neutral swellable polymers,
     which releases the active ingredient in a predetd. manner. For example, a sustained-release tablet composition contained cephalexin 795.32 mg, lactose
     107.68 mg, xanthan gum 31.5 mg, Eudragit NE 30D 52.5 mg, HPMC E5 52.5 mg,
     and magnesium stearate 10.5 mg. Cephalexin release from the tablets was 25.21%, 50.84%, 73.18%, and 84.17% after 1h, 6 h, 10 h and 14 h, resp.
     cephalosporin polymer mixt sustained release tablet
     Dissolution
     Drug bioavailability
     Granulation
         (sustained-release composition of cephalosporin antibiotic containing mixture of
         polymers)
     Polymers, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
         (sustained-release composition of cephalosporin antibiotic containing mixture of
         polymers)
IT
     Drug delivery systems
         (sustained-release; sustained-release composition of cephalosporin
         antibiotic containing mixture of polymers)
     Drug delivery systems
IT
         (tablets, sustained-release; sustained-release composition of cephalosporin
         antibiotic containing mixture of polymers)
                                                             9010-88-2, Eudragit NE
     9000-30-0, Guar gum 9000-40-2, Locust bean gum
     30D 11078-30-1, Galactomannan 11138-66-2, Xanthan gum
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
         (sustained-release composition of cephalosporin antibiotic containing mixture of
         polymers)
                               11111-12-9, Cephalosporin
                                                             15686-71-2, Cephalexin
IT
     153-61-7, Cephalothin
     21593-23-7, Cefaprin
50370-12-2, Cefadroxil
                                                          35607-66-0, Cefoxitin
                              34444-01-4, Cefamandole
                                53994-73-3, Cefaclor
                                                          61270-58-4, Cefonicid
                                       68401-81-0, Ceftizoxime
                                                                    87239-81-4.
     64544-07-6, Cefuroxime axetil
                              92665-29-7, Cefprozil
                                                       117467-28-4, Cefditoren
     Cefpodoxime proxetil
     pivoxil
     RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
         (sustained-release composition of cephalosporin antibiotic containing mixture of
         polymers)
=> b wpix
FILE 'WPIX' ENTERED AT 15:12:08 ON 15 APR 2005
COPYRIGHT (C) 2005 THE THOMSON CORPORATION
                               14 APR 2005
                                                  <20050414/UP>
FILE LAST UPDATED:
MOST RECENT DERWENT UPDATE:
                                                   <200524/DW>
                                   200524
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DERWENT WORLD PATENTS INDEX SUBSCRIBER FILE, COVERS 1963 TO DATE

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>>> FOR A COPY OF THE DERWENT WORLD PATENTS INDEX STN USER GUIDE.
    PLEASE VISIT:
 http://www.stn-international.de/training_center/patents/stn_guide.pdf <<<
>>> FOR DETAILS OF THE PATENTS COVERED IN CURRENT UPDATES, SEE
   http://thomsonderwent.com/coverage/latestupdates/
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>>> FOR INFORMATION ON ALL DERWENT WORLD PATENTS INDEX USER
    GUIDES, PLEASE VISIT: http://thomsonderwent.com/support/userguides/
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>>> NEW! FAST-ALERTING ACCESS TO NEWLY-PUBLISHED PATENT DOCUMENTATION NOW AVAILABLE IN DERWENT WORLD PATENTS INDEX
    FIRST VIEW - FILE WPIFV.
    FOR FURTHER DETAILS: http://www.thomsonderwent.com/dwpifv <<<
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http://thomsonderwent.com/support/dwpiref/reftools/classification/code-revision/
    FOR DETAILS. <<<
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     ANSWER 1 OF 1 WPIX COPYRIGHT 2005 THE THOMSON CORP on STN
     2004-257150 [24]
     2004-257149 [24]
CR
     C2004-100402
     Sustained release composition useful for treating bacterial and microbial
     infection comprises cephalosporin antibiotic, and a mixture of polymers
     containing galactomannans and neutral swellable polymers.
DC
     A96 B02 B07
     BOLDHANE, S P; JINDAL, K C; KSHIRSAGAR, R S (ORCH-N) ORCHID HEALTH CARE; (ORCH-N) ORCHID CHEM & PHARM LTD; (ORCH-N)
IN
     ORCHID HEALTH CARE DIV ORCHID CHEM & PHA
CYC
     103
                      A1 20040226 (200424)* EN
PΙ
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                                                    55
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             DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR
             KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PH PL
             PT RO RU SC SD SE SG SK SL TJ TM TN TR TT TZ UA UG US UZ VC VN YU
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A1 20040520 (200434)
A1 20040303 (200457)
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     US 2004096496
                                                           A61K031-545
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                                                           A61K009-22
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     20020819; US 2004096496 A1 CIP of US 2002-222930 20020819, US 2003-642194
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PRAI WO 2002-IB3320
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     ICM (A61K009-22) A61K031-545
ICS A61K009-20; A61K031-546; A61K031-736
     W02004016251 A UPAB: 20040907
     NOVELTY - A composition (C1) comprises cephalosporin antibiotic, a mixture
     of polymers containing galactomannans and neutral swellable polymers, and
     an excipient.
           DETAILED DESCRIPTION - An INDEPENDENT CLAIM is included for the
     preparation of (C1) involving:
           (1) mixing the active ingredient, excipients and galactomannans in a
           (2) optionally compacting the mixture and sizing it by passing
     through sieve;
           (3) granulating the mixture with neutral swellable polymer;
           (4) drying the granules by either tray drier or fluid bed drier;
           (5) milling the dried granules followed by addition and blending of
     dry binder and lubricant(s);
```

(6) compressing the lubricated granules into tablets using a tablet press; and (7) optionally coating the tablets. ACTIVITY - Antibacterial; Antimicrobial. MECHANISM OF ACTION - Microbial growth inhibitor. USE - For the treatment of bacterial and microbial infections. ADVANTAGE - The composition reduces the administration frequency by maintaining a constant plasma concentration of drug over an extended period of time to ensure sustained effect of active ingredient. The release of active ingredient from the delivery system is controlled by the specific polymers present in the matrix and in specific concentrations, thus allowing blood levels above MIC over extended period of time such that the frequency of the dosage form is released at a rate suitable for once or twice daily administration. Dwg. 0/5 CPĬ AB; DCN CPI: A03-A00A; A12-V01; B02-C02; B04-B01C1; B04-C02A; B04-C02B; B04-C02D; B04-C03; B04-N02; B05-B02C; B07-A02; B10-A07; B10-C04E; B12-M10A;

=> b home

FILE 'HOME' ENTERED AT 15:12:12 ON 15 APR 2005

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FS

FA

MC

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=> d his full
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               3 SEA ABB=ON PLU=ON (L1 OR L2)
L4
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     FILE 'HCAPLUS' ENTERED AT 15:10:26 ON 15 APR 2005
L5
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                E E 3+ALL
                E SWELLING AGENTS/CT
                 E E3+ALL
L12
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                 E POLYMERS/CT
                 E E3+OLD, NT1
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L13
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                E E6+ALL
                E E2+ALL
L14
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L15
                D SCA
                D IDE L15
L16
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L17
                AN? OR MEYPROID? OR MOLVENIN? OR SUNFIBER? OR SUN (W) FIBER?
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JINDAL KOUR CHAND"/AU)

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                    SEL AN 1-3 L41
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L43
                12 SEA ABB=ON PLU=ON L36 OR L42
1.44
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STRUCTURE FILE UPDATES: 14 APR 2005 HIGHEST RN 848555-82-8 DICTIONARY FILE UPDATES: 14 APR 2005 HIGHEST RN 848555-82-8

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 18, 2005

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Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

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ANSWER 1 OF 1 REGISTRY COPYRIGHT 2005 ACS on STN
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RN
     Entered STN: 16 Nov 1984
    Cephalosporin (9CI) (CA INDEX NAME)
CN
OTHER NAMES:
CN
    Ceflin
CN
    Cephalosporins
CN
    Cephalotinum
MF
    Unspecified
    COM, MAN
CI
     STN Files:
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IN Files: ADISNEWS, AGRICOLA, ANABSTR, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CBNB, CEN, CHEMLIST, CIN, CSCHEM, CSNB, DIOGENES, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSCOSEARCH, MEDLINE, NAPRALERT, PIRA, PROMT, TOXCENTER, USPATTULL

EINECS\*\* Other Sources:

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

#### \*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4817 REFERENCES IN FILE CA (1907 TO DATE) 862 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 4821 REFERENCES IN FILE CAPLUS (1907 TO DATE)

#### => d ide 115 tot

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L15 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2005 ACS on STN
      11078-30-1 REGISTRY
      Entered STN: 16 Nov 1984
      D-Galacto-D-mannan (9CI) (CA INDEX NAME)
OTHER NAMES:
      Daikol U 2
CN
CN
      Faibaron S
CN
      Galactomannan
CN
      Guapack PM 1
CN
      Mannan, galacto
      Mannogalactan
CN
      Meyproid 7700
CN
CN
      Molvenin 848
CN
      Sunfiber R
DR
      74505-30-9, 76688-81-8
MF
      Unspecified
      PMS, COM, MAN
CI
      Manual registration
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      STN Files:
      EMBASE, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MSDS-OHS, NAPRALERT, PIRA, PROMT, TOXCENTER, TULSA, USPAT2, USPATFULL
Other Sources: EINECS**, NDSL**, TSCA**
            (**Enter CHEMLIST File for up-to-date regulatory information)
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## \*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1963 REFERENCES IN FILE CA (1907 TO DATE) 263 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 1963 REFERENCES IN FILE CAPLUS (1907 TO DATE)

Qazi 10/642194

Page 4

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FILE COVERS 1907 - 15 Apr 2005 VOL 142 ISS 17 FILE LAST UPDATED: 14 Apr 2005 (20050414/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN
L43
     2004:412577 HCAPLUS
     140:395542
ED
     Entered STN: 21 May 2004
     Sustained release pharmaceutical composition of a cephalosporin
     antibiotic
     Kshirsagar, Rajesh Suresh; Boldhane, Sanjay Parbhatrao
     ; Jindal, Kour Chand
     Orchid Chemicals & Pharmaceuticals Limited,
     India
S0
     U.S. Pat. Appl. Publ., 21 pp., Cont.-in-part of U.S. Ser. No. 222,930.
     CODEN: USXXCO
DT
     Patent
     English
LA
     ICM A61K031-545
IC
         A61K009-20; A61K031-736
     ICS
     424465000; 514200000; 514054000
NCL
CC
     63-6 (Pharmaceuticals)
FAN. CNT 3
     PATENT NO.
                          KIND
                                  DATE
                                              APPLICATION NO.
                                                                       DATE
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                                  20040520
                                              US 2003-642194
                                                                       20030818
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     US 2004033262
                                  20040219
                                              US 2002-222930
                                                                       20020819
                           A1
     WO 2004016250
                                  20040226
                                               WO 2002-IB3320
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                                       DM, DZ,
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                         A61K009-20; A61K031-736
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NCL
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                        A61K031/736
     Sustained release pharmaceutical compns. comprising at least a
     cephalosporin antibiotic, a mixture of polymers comprising of
     galactomannans and neutral swellable polymers, and other
     pharmaceutically acceptable excipients are described. The composition
     comprises about 30% to about 90% by weight of a cephalosporin
     antibiotic; about 1% to about 30% by weight of said mixture of polymers
     comprising from about 0.1% to about 15% by weight of galactomannans
     , and about 0.1% to about 15% of neutral swellable polymer by weight of
     sustained release composition
     cephalosproin antibiotic sustained release tablet .
     Drug delivery systems
        (carriers; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IΤ
     Lubricants
        (pharmaceutical; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
     Swelling agents
        (polymers; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
     Antibiotics
     Binders
     Drying
     Mixing
     Plasticizers
     Sieving
        (sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
    Gelatins, biological studies
RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
IT
     (Biological study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
     Polyesters, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
IT
    Drug delivery systems
        (sustained-release; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
    Polymers, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (swellable; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
    Drug delivery systems
        (tablets, coated; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
     Drug delivery systems
        (tablets; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
     Fats and Glyceridic oils, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
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        (vegetable, hydrogenated; sustained-release pharmaceutical composition of a cephalosporin antibiotic)
    7631-86-9, Silicon dioxide, biological studies
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     (Biological study); USES (Uses)
        (colloidal; sustained-release pharmaceutical composition of a
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    9004-34-6, Cellulose, biological studies
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     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
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9005-25-8, Starch, biological studies RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (pregelatinized; sustained-release pharmaceutical composition of a cephalosporin antibiotic) 9003-39-8, Polyvinylpyrrolidone 9004-62-0, Hydroxyethylcellulose 9004-64-2, Hydroxypropyl cellulose 9004-65-3, Hydroxypropyl IT methylcellulose RL: MOA (Modifier or additive use); POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (sustained-release pharmaceutical composition of a cephalosporin antibiotic) 57-11-4, Stearic acid, biological studies 57-50-1, Sucrose, biological studies 557-04-0, Magnesium stearate 557-05-1, Zinc stearate 1592-23-0, Calcium stearate 9000-30-0, Guar gum 11138-66-2, Xanthan gum 14807-96-6, Talc, biological studies RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (sustained-release pharmaceutical composition of a cephalosporin antibiotic) 153-61-7, Cephalothin 11078-30-1, Galactomannan ΙT 11111-12-9D, Cephalosporins, derivs. 15686-71-2, Cephalexin 21593-23-7, Cephapirin **34444-01-4**, Cefamandole 35607-66-0, Cefoxitin **50370-12-2** , Cefadroxil **53994-73-3**, Cefaclor 61270-58-4, Cefonicid 64544-07-6, Cefuroxime axetil **68401-81-0**, Ceftizoxime 87239-81-4, Cefpodoxime proxetil 92665-29-7, Cefprozil 117467-28-4. Cefditoren pivoxil RL: PEP (Physical, engineering or chemical process); PYP (Physical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses) (sustained-release pharmaceutical composition of a cephalosporin antibiotic) 9004-32-4 9010-88-2, Ethyl acrylate-methyl methacrylate copolymer IT 9050-04-8, Calcium carboxymethyl cellulose 25086-89-9, Plasdone S-630 RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (sustained-release pharmaceutical composition of a cephalosporin antibiotic) 50-70-4, Sorbitol, biological studies 50-99-7, Glucose, biological studies 63-42-3, Lactose 69-65-8, Mannitol 69-79-4, Maltose ΙT 9004-53-9, Dextrin 66828-18-0, Dextrate 74811-65-7, Croscarmellose sodium RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (sustained-release pharmaceutical composition of a cephalosporin antibiotic) 153-61-7, Cephalothin RL: PEP (Physical, engineering or chemical process); PYP (Physical IT process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); (sustained-release pharmaceutical composition of a cephalosporin antibiotic) RN 153-61-7 HCAPLUS 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R, 7R)-(9CI)(CA INDEX NAME)

Absolute stereochemistry.

L43 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN

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     140:205146
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     Entered STN: 29 Feb 2004
     Sustained release pharmaceutical compositions of a cephalosporin
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       Jindal, Kour Chand
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     Chemicals & Pharmaceuticals Ltd., India
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     PCT Int. Appl., 55 pp.
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                          A61K009/20H6B; A61K009/20H6F; A61K031/545; A61K031/546;
                          A61K031/736
     Sustained release pharmaceutical compns. comprising at least a
AB
     cephalosporin, a mixture of polymers comprising of
     galactomannans and neutral swellable polymers, and other
     pharmaceutically acceptable excipients are described. The composition
     comprises 30-90% of a cephalosporin antibiotic, 1-30% mixture of
     polymers comprising 0.1-15% galactomannans, and 0.1-15% by weight
     of a neutral swellable polymer. Thus, tablets contained cephalexin monohydrate 75.73, lactose 6.27, xanthan gum 7.0, Eudragit NE 30D 7.0,
     HPMC E5 31.5, Mg stearate 1.0 and water qs to 100%.
     sustained release cephalosporin polymer
     Drug delivery systems
         (granules, sustained release; sustained release pharmaceutical compns.
         of cephalosporins)
IT
     Binders
     Dissolution
     Drug bioavailability
     Human
     Lubricants
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Page 8

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(sustained release pharmaceutical compns. of cephalosporins)
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     Gelatins, biological studies
     Polymers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained release pharmaceutical compns. of cephalosporins)
IT
     Drug delivery systems
         (sustained-release; sustained release pharmaceutical compns. of
         cephalosporins)
ΙT
     Drug delivery systems
         (tablets, sustained-release; sustained release pharmaceutical compns.
         of cephalosporins)
     Fats and Glyceridic oils, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (vegetable, hydrogenated; sustained release pharmaceutical compns. of
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(1) Lupin Lab Ltd; WO 0241876 A 2002 HCAPLUS
   Shankar, M; WO 02067943 A 2002 HCAPLUS
(3) Squibb Bristol Myers Co; WO 9846213 A 1998 HCAPLUS
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Absolute stereochemistry.

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     Entered STN: 22 Feb 2004
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     US 2004096496
                                  20040520
                                               US 2003-642194
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PRAI IN 2002-MA601
                                  20020816
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        2002-222930
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     WO 2002-IB3320
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CLASS
 PATENT NO.
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                         PATENT FAMILY CLASSIFICATION CODES
 US 2004033262
                  ICM
                         A61K031-545
                         A61K031-736; A61K009-20; A61K009-22
                  ICS
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 US 2004033262
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                         A61K009/20H6B; A61K009/20H6F; A61K031/545; A61K031/546;
                         A61K031/736
                  ECLA
                         A61K031/545; A61K031/546
 US 2004096496
AB A sustained release pharmaceutical composition comprising at least a
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cephalosporin antibiotic and a mixture of polymers and other
     pharmaceutically acceptable excipients is described. Polymers are
     selected from mixture of galactomannans and neutral swellable
     polymers, which releases the active ingredient in a predetd. manner. For
     example, a sustained-release tablet composition contained cephalexin 795.32 mg,
     lactose 107.68 mg, xanthan gum 31.5 mg, Eudragit NE 30D 52.5 mg, HPMC E5
     52.5 mg, and magnesium stearate 10.5 mg. Cephalexin release from the tablets was 25.21%, 50.84%, 73.18%, and 84.17% after 1h, 6 h, 10 h and 14
     cephalosporin polymer mixt sustained release tablet
     Dissolution
     Drug bioavailability
     Granulation
        (sustained-release composition of cephalosporin antibiotic containing
        mixture of polymers)
     Polymers, biological studies
RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (sustained-release composition of cephalosporin antibiotic containing
        mixture of polymers)
IT
     Drug delivery systems
        (sustained-release; sustained-release composition of cephalosporin
        antibiotic containing mixture of polymers)
IT
     Drug delivery systems
         (tablets, sustained-release; sustained-release composition of
        cephalosporin antibiotic containing mixture of polymers)
IT
     9000-30-0, Guar gum 9000-40-2, Locust bean gum 9010-88-2, Eudragit NE
     30D 11078-30-1, Galactomannan 11138-66-2, Xanthan
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (sustained-release composition of cephalosporin antibiotic containing
        mixture of polymers)
     153-61-7, Cephalothin 11111-12-9, Cephalosporin
ΙT
     15686-71-2, Cephalexin 21593-23-7, Cefaprin
                                35607-66-0, Cefoxitin 50370-12-2
     34444-01-4, Cefamandole
       Cefadroxil 53994-73-3, Cefaclor 61270-58-4, Cefonicid
     64544-07-6, Cefuroxime axetil 68401-81-0, Ceftizoxime
     87239-81-4, Cefpodoxime proxetil 92665-29-7, Cefprozil
                                                                   117467-28-4,
     Cefditoren pivoxil
     RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (sustained-release composition of cephalosporin antibiotic containing
        mixture of polymers)
    11078-30-1, Galactomannan
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (sustained-release composition of cephalosporin antibiotic containing
        mixture of polymers)
RN
     11078-30-1 HCAPLUS
     D-Galacto-D-mannan (9CI)
                                (CA INDEX NAME)
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L44 ANSWER 1 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN
     2004:267168 HCAPLUS
AN
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     140:309365
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     Entered STN:
                   01 Apr 2004
     Surface dissolution and/or bulk erosion controlled release pharmaceutical
     compositions
     Shefer, Adi; Shefer, Samuel David
IN
PA
S<sub>0</sub>
     U.S. Pat. Appl. Publ., 21 pp.
     CODEN: USXXCO
DT
     Patent
     English
     ICM A61K009-00
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ICS A61K009-127
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                                              APPLICATION NO.
                                                                      DATE
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PRAI US 2002-255289
                                  20020926
CLASS
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                  CLASS
                         PATENT FAMILY CLASSIFICATION CODES
 US 2004062778
                  ICM
                         A61K009-00
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                         A61K009-127
                         424400000
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AB
     The present invention relates to a controlled release system comprising
     matrix compns. which control the lag time and release rate of the composition,
     as well as pharmaceutical and other active ingredients included in the
     composition, through surface dissoln. and/or bulk erosion of the system.
     controlled release system can be used to target and control the release of
     active ingredients onto certain regions of the gastrointestinal tract
     including the stomach and the small intestine. The matrix compns. of the
     present invention can be comprised of the following components: a wax
     material, fat material, water sensitive material and surface active
     material. Thus, a composition contained carnauba wax 30.0, beeswax 5.0,
     glyceryl monostearate 10.0. Tween-20 15.0, lactitol 30.0, and BaSO4 10.0%.
     controlled release pharmaceutical bulk erosion
     Fats and Glyceridic oils, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
IT
         (Japan wax; surface dissoln. and/or bulk erosion controlled release
        pharmaceutical compns.)
IT
     Imaging agents
        (NMR contrast; surface dissoln. and/or bulk erosion controlled release
        pharmaceutical compns.)
IT
     Imaging agents
        (acoustic imaging contrast agents; surface dissoln. and/or bulk erosion
        controlled release pharmaceutical compns.)
IT
     Positron-emission tomography
     Tomography
        (agents for; surface dissoln. and/or bulk erosion controlled release
        pharmaceutical compns.)
IT
     Carboxylic acids, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (alkyl esters; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.)
IT
     Quaternary ammonium compounds, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (alkylbenzyldimethyl, chlorides; surface dissoln. and/or bulk erosion
        controlled release pharmaceutical compns.)
     Surfactants
IT
        (amphoteric; surface dissoln. and/or bulk erosion controlled release
        pharmaceutical compns.)
IT
     Heart, disease
        (angina pectoris; surface dissoln. and/or bulk erosion controlled
        release pharmaceutical compns.)
IT
     Waxes
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (animal; surface dissoln. and/or bulk erosion controlled release
        pharmaceutical compns.)
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(anionic; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Heart, disease (arrhythmia; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Infection (bacterial; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Prostate gland, disease IT (benign hyperplasia; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Hyperplasia (benign prostatic; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Surfactants (cationic; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Inflammation Intestine, disease (colitis; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Intestine (colon; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Nose, disease (congestion; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Drug delivery systems (controlled-release; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Monoglycerides ΙT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (cottonseed-oil, hydrogenated; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) ΙT Mental disorder (depression; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Castor oil RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (derivs.; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Diglycerides RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (digalactosyl; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Waxes RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (emulsifying; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Fatty acids, biological studies IT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (essential; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Fatty acids, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (esters; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Alcohols, biological studies Amides, biological studies Esters, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (ethoxylated; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) ΙT Heart, disease (failure; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Vein, disease

(hemorrhoid; surface dissoln. and/or bulk erosion controlled release

pharmaceutical compns.)

IT Castor oil RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hydrogenated, ethoxylated; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) ΙT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hydrogenated; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Syrups (sweetening agents) (hydrolyzed starch; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Dental materials and appliances Prosthetic materials and Prosthetics (implants; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Sexual behavior (impotence; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Thyroid gland (inhibitors; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Alcohols, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (lanolin; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) ΙT Hydrocarbon waxes, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (microcryst.; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Drug delivery systems (microspheres; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Antibodies and Immunoglobulins RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (monoclonal; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) ΙT Drug delivery systems (nanospheres; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Surfactants (nonionic; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Anti-inflammatory agents (nonsteroidal; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) Alcohols, biological studies · IT Esters, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (propoxylated; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) ΙT Infection (protozoal; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Intestine (small; surface dissoln, and/or bulk erosion controlled release pharmaceutical compns.) IT Muscle, disease (spasm; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Brain, disease (stroke; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT Analgesics Anthelmintics Anti-inflammatory agents Antianginal agents Antiarrhythmics Antibacterial agents Anticoagulants Anticonvulsants

Antidepressants Antidiabetic agents Antiemetics Antihistamines Antihypotensives Antimalarials Antimigraine agents Antiobesity agents Antiparkinsonian agents Antitussives Antiviral agents Anxiety Anxiolytics Appetite depressants Beeswax Blood coagulation Bronchodilators Cognition Cognition enhancers Contraceptives Cosmetics Cough Decongestants Diabetes mellitus Digestive tract Dissolution Diuresis Diuretics Epilepsy Expectorants Fungicides Gout Granulation Gums and Mucilages Hydrocolloids Hydrophile-lipophile balance value Hypertension Hypnotics and Sedatives Hypotension Imaging agents Immunomodulators Immunosuppressants Immunosuppression Inflammation Inotropics Laxatives Leukotriene antagonists Malaria Medical goods Mental disorder Muscarinic antagonists Muscle relaxants Mycosis Neoplasm Nutrition, animal Osteoporosis Ozocerite Pain Parkinson's disease Particle size distribution Protozoacides Psychotropics Sleep Spheronization Stomach Surfactants

Tranquilizers Vaccines Vasoconstriction Vasoconstrictors

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Vasodilation
     Vasodilators
     Vomiting
         (surface dissoln. and/or bulk erosion controlled release pharmaceutical
        compns.)
     Amine oxides
     Candelilla wax
     Carbohydrates, biological studies
     Cardiolipins
     Carnauba wax
     Corticosteroids, biological studies
     Cytokines
     Diglycerides
     Fats and Glyceridic oils, biological studies
     Fatty acids, biological studies
     Gelatins, biological studies
     Glutens
     Glycerides, biological studies
     Hormones, animal, biological studies
     Hydrocarbon waxes, biological studies
     Interferons
     Isoprenoids
     Keratins
     Lecithins
     Lipids, biological studies
     Macrolides
     Monoglycerides
     Oligomers
     Oligonucleotides
     Oligosaccharides, biological studies
     Opioids
     Peptides, biological studies
     Phosphatidic acids
     Phosphatidylcholines, biological studies
     Phosphatidylethanolamines, biological studies
     Phosphatidylgly cerols\\
     Phosphatidylinositols
     Phosphatidylserines
     Phospholipids, biological studies
     Polyamides, biological studies
Polyesters, biological studies
     Polymers, biological studies
     Polyoxyalkylenes, biological studies
Polysaccharides, biological studies
     Prostaglandins
     Proteins
     Sex hormones
     Shellac
     Sphingolipids
     Steroids, biological studies
     Terpenes, biological studies
     Tocopherols
     Tumor necrosis factors
     Vitamins
     Waxes
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (surface dissoln. and/or bulk erosion controlled release pharmaceutical
        compns.)
ΙT
     Drug delivery systems
         (tablets, controlled-release; surface dissoln. and/or bulk erosion
        controlled release pharmaceutical compns.)
     Fats and Glyceridic oils, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
IT
         (vegetable, hydrogenated; surface dissoln. and/or bulk erosion
        controlled release pharmaceutical compns.)
ΙT
     Infection
         (viral; surface dissoln. and/or bulk erosion controlled release
        pharmaceutical compns.)
IT
     Surfactants
         (zwitterionic; surface dissoln. and/or bulk erosion controlled release
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pharmaceutical compns.) IT Adrenoceptor antagonists (β-; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (β-; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) 9015-94-5, Renin, biological studies IΤ RL: BSU (Biological study, unclassified); BIOL (Biological study) (inhibitor; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) 9001-92-7, Protease IT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (inhibitor; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) IT 329900-75-6, COX-2 RL: BSU (Biological study, unclassified); BIOL (Biological study) (inhibitors; surface dissoln. and/or bulk erosion controlled release pharmaceutical compns.) 50-69-1, Ribose 50-70-4, Sorbitol, biological studies 50-78-2, As 50-99-7, Glucose, biological studies 50-99-7D, Glucose, esters 56-86-0D, Glutamic acid, acyl derivs. 57-48-7, Fructose, biological IT 50-78-2, Aspirin 57-50-1, Sucrose, biological studies 57-50-1D, Sucrose, esters 57-55-6D, Propylene glycol, esters 57-88-5, Cholesterol, biological studies 58-86-6, Xylose, biological studies 59-23-4, Galactose, biological studies 69-65-8, Mannitol 69-72-7D, derivs. 69-79-4, Maltose 79-10-7D, Acrylic acid, polymers 79-41-4D, Methacrylic acid, 87-99-0, Xylitol 88-99-3D, Phthalic acid, derivs. esters, polymers 107-21-1D, Ethylene glycol, esters 107-97-1D, 89-78-1, Menthol Sarcosinic acid, acyl derivs. 111-46-6D, Diethylene glycol, esters 112-80-1, Oleic acid, biological studies 124-30-1, Stearylamine 143-07-7, Lauric acid, biological studies 151-21-3, Sodium lauryl sulfate, biological studies 373-49-9, Palmitoleic acid 544-64-9, Myristoleic acid 544-66-1, Physeteric acid 555-43-1, Glycerol 593-39-5, 585-86-4, Lactitol 585-88-6, Maltitol tristearate Petroselinic acid 1256-86-6, Cholesterol sulfate 1510-21-0, Cholesterol hemisuccinate 2462-63-7, Dioleoylphosphatidylethanolamine 2644-64-6, Dipalmitoylphosphatidylcholine 3458-28-4, Mannose 4539-70-2, Distearoylphosphatidylcholine 7664-93-9D, Sulfuric acid, 4539-70-2, Distearoylphosphatidylcholine esters 7727-43-7, Barium sulfate 7778-18-9, Calcium sulfate 8050-81-5, Simethicone 9000-01-5, Acacia gum 9000-65-1, Tragacanth 9002-72-6, Growth hormone 9002-79-3, Melanocyte-stimulating hormone 9002-89-5, Polyvinyl alcohol 9003-01-4D, Polyacrylic acid, derivs. 9003-05-8, Polyacrylamide 9003-39-8, Polyvinylpyrrolidone 9004-1 Insulin, biological studies 9004-32-4, Carboxymethyl cellulose 9004-10-8. 9004-34-6, Cellulose, biological studies 9004-34-6D, Cellulose, derivs. 9004-38-0, Cellulose acetate phthalate 9004-62-0, Hydroxyethyl cellulose 9005-25-8, Starch, biological studies 9005-25-8D, Starch, derivs. 9005-64-5, Tween 20 9006-26-2, Ethylene-maleic anhydride copolymer 9007-12-9, Calcitonin 9011-16-9, Maleic anhydride-methylvinyl ether copolymer 9026-81-7, Nuclease 9032-50-2, Methyl cellulose phthalate 9050-31-1, Hydroxypropyl methyl cellulose phthalate 9050-36-6, Maltrin M 180 9054-89-1, Superoxide dismutase 9062-05-9, Hydroxypropyl carboxymethyl cellulose 9083-87-8 11078-30-1,
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Cephalosporin 11138-66-2, Xanthan 12441-09-7D, Sorbitan, 15687-27-1, Ibuprofen 18656-38-7, Dimyristoylphosphatidylcholin e 18656-40-1, Dilauroylphosphatidylcholine 24529-88-2 25322-68-3, Polyethylene oxide 25322-69-4 25618-55-7D, Polyglycerol, esters with fatty acids 29385-00-0, Isolauric acid 30399-84-9, Isostearic acid 31566-31-1, Glyceryl monostearate 32844-67-0, Isopalmitic acid 37324-30-4, Hydroxypropyl cellulose phthalate 37353-59-6, Hydroxymethyl cellulose 51110-01-1, Somatostatin 52907-01-4, Cellulose acetate trimellitate 53237-50-6 53714-56-0, Leuprolide 56509-23-0, Sodium cellulose acetate phthalate 59865-13-3, Cyclosporin 62571-86-2, 62683-29-8, Colony stimulating factor 64519-82-0, Palatinit Captopril 64792-89-8, Dibehenoylphosphatidylcholine 65437-21-0, Isomyristic acid 66829-29-6, Hi-Cap 100 67896-63-3, Dipentadecanoylphosphatidylcholine 68354-99-4 68424-04-4, Polydextrose 68737-67-7,

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Dioleoylphosphatidylcholine
                                     71138-97-1, Hydroxypropylmethyl cellulose
     acetate succinate 71259-34-2
                                        76822-97-4
                                                       78543-25-6,
     1-Hexadecyl-2-palmitoylglycerophosphoethanolamine
                                                             82196-85-8
     83172-32-1, Ditricosanoylphosphatidylcholine
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                    106392-12-5, Poloxamer
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     Ondansetron
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                                   143475-87-0
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     129385-16-6
                                           252856-84-1, Polyvinyl acetate hydrogen
     Dilignoceroylphosphatidylcholine
     phthalate 676259-03-3 676270-47-6
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     11078-30-1, Galactomannan 11111-12-9,
ΙT
     Cephalosporin
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (surface dissoln. and/or bulk erosion controlled release pharmaceutical
         compns.)
RN
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                            (CA INDEX NAME)
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     Entered STN: 26 Dec 2003
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TI
     Film-shaped mucoadhesive administration form for administering cannabis
     active ingredients
IN
     Wessling, Werner
     LTS Lohmann Therapie-Systeme A.-G., Germany
PA
S<sub>0</sub>
     PCT Int. Appl., 12 pp.
     CODEN: PIXXD2
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PRAI DE 2002-10226494 A 20020614
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CLASS
 PATENT NO.
                  CLASS
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 WO 2003105800
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                          A61K009-70; A61K035-78
A61K009/00M18D; A61K009/70B
A61K009/00M18D; A61K009/70B
                  ICS
 WO 2003105800
                  ECLA
 DE 10226494
                  ECLA
     The invention relates to a film-shaped, mucoadhesive administration form
     containing at least one active ingredient from the group of cannabis active
     ingredients. Sublingual or buccal films are prepared that also contain
     swellable polymers, flavoring substances, fillers, dyes, emulsifiers,
     softeners, permeation enhancers, antioxidants, buffers, and preservatives.
     Active ingredient content is 0.1-20 weight/weight%; preferably 0.5-10 %; the
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thickness of the films is 0.05-0.5 mm. The compns. are applied as pain
     killers for cancer and AIDS patients and to alleviate other diseases.
     cannabis tetrahydrocannabinol drug mucoadhesive film buccal sublingual
ΙT
     Drug delivery systems
        (buccal; film-shaped mucoadhesive administration form for administering
        cannabis active ingredients)
ΙT
     Essential oils
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (chamomile; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
    Mental disorder
        (dementia; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
     Opioids
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
        (dependence; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
    Nervous system, disease (dystonia; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
    Essential oils
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (eucalyptus; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
     AIDS (disease)
IT
     Alcoholism
     Alzheimer's disease
     Analgesics
     Anorexia
     Anticonvulsants
     Antiemetics
     Antimigraine agents
     Antioxidants
     Arthritis
     Asthma
     Buffers
     Cachexia
     Chemotherapy
     Drug dependence
     Drug withdrawal
     Dyes
     Dysmenorrhea
     Emulsifying agents
     Epilepsy
     Fillers
     Flavor
     Glaucoma (disease)
     Hepatitis
     Human
     Multiple sclerosis
     Neoplasm
     Parkinson's disease
     Permeation enhancers
     Plasticizers
     Preservatives
     Sweetening agents
     Swelling, physical
        (film-shaped mucoadhesive administration form for administering
        cannabis active ingredients)
     Collagens, biological studies
     Gelatins, biological studies
     Polyoxyalkylenes, biological studies
     Turpentine oil
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (film-shaped mucoadhesive administration form for administering
        cannabis active ingredients)
IT
     Drug delivery systems
        (films; film-shaped mucoadhesive administration form for administering
        cannabis active ingredients)
     Cannabis sativa
IT
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(marijuana, exts. or oils of; film-shaped mucoadhesive administration
        form for administering cannabis active ingredients)
IT
     Headache
         (migraine; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
     Pain
         (neuropathic; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
         (of drug film; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
     Drug delivery systems
         (sublingual; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
     Polymers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (swellable; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
     Essential oils
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (thyme, Thymus vulgaris; film-shaped mucoadhesive administration form
        for administering cannabis active ingredients)
IT
     In jury
         (trauma; film-shaped mucoadhesive administration form for administering
        cannabis active ingredients)
ΙT
     Disease, animal
         (wasting; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
IT
     12794-10-4. Benzodiazepine
     RL: ADV (Adverse effect, including toxicity); BIOL (Biological study)
         (dependence; film-shaped mucoadhesive administration form for
        administering cannabis active ingredients)
     60-12-8, Phenylethanol 79-10-7D, Acrylic acid, esters, polymers
     79-92-5, Camphene 89-78-1, Menthol
                                               92-48-8, 6-Methyl coumarin
     106-22-9, Citronellol 119-36-8, Methylsalicylate
                                                               121-32-4,
     Ethylvanillin 123-86-4, Acetic acid-n-butyl ester
                                                                 138-86-3, Limonene
     470-82-6, Eucalyptol 521-35-7, Cannabinol 1330-16-1, Pine 1972-08-3, Tetrahydrocannabinol 6066-49-5, Butylphthalide
                                                        1330-16-1, Pinene
                        9000-65-1, Tragacanth gum
9003-01-4, Polyacrylic acid
     Carrageenan gum
                                                       9000-69-5, Pectins
     9002-18-0, Agar
                                                          9003-05-8, Polyacrylamide
     9003-39-8, Polyvinylpyrrolidone 9004-32-4, Carboxymethyl cellulose
     9004-57-3, Ethyl cellulose 9004-62-0, Hydroxyethyl cellulose
     9004-64-2, Hydroxypropyl cellulose 9005-18-9, Propyl cellulose 9005-25-8, Starch, biological studies 9005-25-8D, Starch, derivs. 9005-32-7, Alginic acid 9012-36-6, Agarose 9012-76-4, Chitosan
     9036-66-2, Arabinogalactan 9057-02-7, Pullulan 11078-30-1, Galactomannan 13956-29-1, Cannabidiol 20675-51-8, Cannabichromene 25322-68-3, Polyethylene oxide 37353-59-6
                                                              37353-59-6.
     Hydroxymethyl cellulose
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (film-shaped mucoadhesive administration form for administering
        cannabis active ingredients)
IT
     11078-30-1, Galactomannan
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (film-shaped mucoadhesive administration form for administering
        cannabis active ingredients)
     11078-30-1 HCAPLUS
     D-Galacto-D-mannan (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     ANSWER 3 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN
L44
     2003:1215 HCAPLUS
AN
DN
     138:61315
     Entered STN: 02 Jan 2003
ED
     Controlled and sustained release dosage forms containing hydrophilic
     carriers and diffusion enhancers
IN
     Chhabra, Harinderpal; Sarkar, Shyamal K.
PA
     USA
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S<sub>0</sub>

U.S., 23 pp.

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CODEN: USXXAM
DT
      Patent.
      English
IC
      ICM A61K009-22
      ICS
          A61K009-24; A61K009-30
     424474000; 424468000; 424470000; 424472000; 424475000; 514770000; 514772300; 514777000; 514778000; 514779000
     63-6 (Pharmaceuticals)
FAN. CNT 1
      PATENT NO.
                                                                                  DATE
                              KIND
                                      DATE
                                                     APPLICATION NO.
                               B1
                                                     US 1999-358732
                                                                                  19990721
     US 6500459
                                       20021231
                                                                                  20000721
                                       20010121
                                                     CA 2000-2314298
      CA 2314298
                               AA
PRAI US 1999-358732
                                       19990721
                               Α
CLASS
 PATENT NO.
                    CLASS PATENT FAMILY CLASSIFICATION CODES
 US 6500459
                    ICM
                             A61K009-22
                             A61K009-24; A61K009-30
                    ICS
                            424474000; 424468000; 424470000; 424472000; 424475000; 514770000; 514772300; 514777000; 514778000; 514779000 A61K009/00L4; A61K009/28K
                    NCL
                    ECLA
 US 6500459
     A pharmaceutical composition for controlled onset and sustained release of an
      active ingredient, comprises: (i) a core comprising: (a) an active
      ingredient; (b) a hydrophilic carrier; (c) a hydrodynamic diffusion
     enhancer; and optionally (d) conventional excipients selected from the group consisting of binders, fillers and lubricants and combinations
      thereof; and (ii) a functional coating membrane surrounding the core.
     Thus, 240 g verapamil-HCl was sieved through a mesh sieve and blended with 150 g E50 premium HPMC. To this blend was added 270.0 g croscarmellose
      sodium and mixed for 15 min. This blend was granulated with PVP K-29/32
     solution in iso-PrOH (30% weight/weight). The wet mass obtained in the above step was dried at 60° for 3 h. After drying, the granules were passed a mesh sieve. The granules were then mixed with 2.5 g of Magnesium Stearate
      and 15 g of Stearic acid in a V blender. This granule blend was
     compressed in a tablet press by using appropriate size tooling. The granules were then mixed with 2.5 g of Mg stearate and 15 g of stearic
      acid in a V blender. This granule blend was compressed in a tablet press
      by using appropriate size tooling. These tablets were then coated by
      using a perforated coating pan. A seal coating membrane was applied on
      the surface of tablets to achieve a weight gain of 1.66% of the weight of the
      core. The seal coating dispersion of Opadry Clear in water at 10% was
      sprayed on to the surface of the tablets by using a perforated coating
      pan.
ST
      controlled release drug hydrophilic carrier diffusion enhancer; sustained
      release drug hydrophilic carrier diffusion enhancer
IT
     Erythropoiesis
          (agents for; controlled and sustained release dosage forms containing
         hydrophilic carriers and diffusion enhancers)
IT
      Anti-infective agents
      Anti-inflammatory agents
      Antianginal agents
      Antiarrhythmics
      Antiasthmatics
      Antibacterial agents
      Anticholesteremic agents
      Anticonvulsants
      Antidiabetic agents
      Antidiarrheals
      Antiemetics
      Antihistamines
      Antihypertensives
      Antitussives
      Anxiolytics
      Appetite depressants
      Binders
      Coating materials
      Decongestants
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Diuretics

Qazi 10/642194

Page 21

Expectorants Hypnotics and Sedatives Hypolipemic agents Laxatives Lubricants Molecular weight distribution Nervous system stimulants Plasticizers Psychotropics Thyroid gland, disease Vasoconstrictors Vasodilators Viscosity (controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) ΙT Vitamins Waxes RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT Drug delivery systems (controlled-release; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT Vasodilators (coronary; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT Clays, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (diffusion enhancer; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT Diffusion (enhancers for; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT Drug delivery systems (granules; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) Polyoxyalkylenes, biological studies IT Polysaccharides, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hydrophilic carrier; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT Protamines RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (sulfates; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) ΙT Drug delivery systems (sustained-release; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT Drug delivery systems (tablets, controlled-release; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) ΙT Drug delivery systems (tablets, sustained-release; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) 9003-39-8D, crosslinked IT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (Crospovidone, diffusion enhancer; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) 50-70-4, Sorbitol, biological studies 50-99-7, Dextrose, biological 57-50-1, Sucrose, biological studies 63-42-3, Lactose studies 69-65-8, Mannitol 9004-34-6, Cellulose, biological studies 9050-36-6, Maltodextrin Starch, derivs. RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (binder; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) IT 50-02-2, Dexamethasone 50-24-8, Prednisolone 50-33-9, Phenylbutazone, 50-34-0, Propantheline bromide 50-47-5, biological studies Desmethylimipramine 50-48-6, Amitriptyline 50-49-7, Imipramine 50-52-2, Thioridazine 50-53-3, Chlorpromazine, biological studies

50-54-4, Quinidine sulfate 50-81-7, Ascorbic acid, biological studies 51-06-9, Procainamide 51-48-9, Thyroxin, biological studies 51-52-5, Propylthiouracil 52-01-7, Spironolactone 52-28-8, Codeine phosphate 52-53-9, Verapamil 52-86-8, Haloperidol 53-03-2, Prednisone 53-86-1, Indomethacin 54-31-9, Furosemide 55-56-1, Chlorhexidine 55-63-0, Glyceryl trinitrate 57-66-9, Probenecid 57-96-5, Sulphinpyrazone 58-55-9, Theophylline, biological studies 58-73-1, Diphenhydramine 58-74-2, Papaverine 58-93-5, Hydrochlorthiazide 59-02-9, α-Tocopherol 59-30-3, FOlic acid, biological studies 59-43-8, Thiamin, biological studies 59-67-6, Nicotinic acid, biological studies 60-13-9, Amphetamine sulphate 60-32-2, ε-Aminocaproic acid 61-75-6, Bretylium tosylate 65-23-6, Pyridoxine 69-23-8, Fluphenazine 76-42-6, Oxycodeinone 77-36-1, Chlorthalidone 78-11-5, Pentaerythritol tetranitrate 86-22-6, Brompheniramine 86-54-4, Hydralazine 87-33-2, Sorbide dinitrate 90-82-4, Pseudoephedrine 93-14-1, Guaiphenesin 94-20-2, Diabenese 113-15-5, Ergotamine 113-45-1, Methylphenidate 113-92-8 114-07-8, Erythromycin 117-89-5, Trifluperazine 123-03-5, Cetylpyridinium chloride 125-71-3, Dextromethorphan 128-62-1, Noscapine 134-80-5, Diethylpropion hydrochloride 146-22-5, Nitrazepam 152-11-4, Verapamil hydrochloride 298-46-4, Carbamazepine 299-28-5, Calcium gluconate 315-30-0, Allopurinol 317-34-0, Aminophylline 364-62-5, Metoclopramide 389-08-2, Nalidixic acid 396-01-0, Triamterene 404-82-0, Fenfluramine hydrochloride 439-14-5, Diazepam 456-59-7, Cyclandelate 480-30-8, Dichloral phenazone 525-66-6, Propranolol 554-13-2, Lithium carbonate 555-30-6, Methyldopa 577-11-7, Dioctyl sodium sulfosuccinate 603-50-9, Bisacodyl 63 Phenytoin sodium 638-23-3, Carbocysteine 846-49-1, Lorazepam 846-50-4, Temazepam 915-30-0, Diphenoxylate 1069-66-5, SOdium valproate 1309-42-8, Magnesium hydroxide 1404-88-2, Tyrothricin 1420-53-7, Codeine sulfate 1622-61-3, Clonazepam 1668-19-5, Doxepine 1847-24-1, Flucloxacillin sodium 2706-50-5, Amphetamine hydrochloride 3200-06-4, Naftidrofuryl oxalate 3737-09-5, Disopyramide 3930-20-9, Sotalol 4205-90-7, Clonidine 5104-49-4, Flurbiprofen 5874-97-5, Orciprenaline sulfate 5965-13-9 6493-05-6, Pentoxifylline Triiodothyronine 7054-25-3, Quinidine gluconate 7683-59-2, Isoproterenol 7720-78-7, Ferrous sulphate 8067-24-1, Co-dergocrine mesylate 9004-10-8, Insulin, biological studies 10238-21-8, Glyburide 10347-81-6, Maprotiline hydrochloride 10377-48-7, Lithium sulfate 14663-23-1, Dantrolene sodium 14838-15-4, Phenylpropanolamine 15307-86-5, Diclofenac 15686-71-2, Cephalexin 15687-27-1, Ibuprofen 17617-23-1, Flurazepam 17693-51-5, Promethazine theoclate 18559-94-9, Salbutamol 21256-18-8, Oxaprozin 21645-51-2, Aluminum hydroxide, biological studies 21829-25-4, Nifedipine 22204-53-1, Naproxen 23031-32-5, Terbutaline sulfate 23887-31-2, Clorazepate 25812-30-0, Gemfibrozil 26652-09-5, Ritodrine 26807-65-8, Indapamide 29094-61-9, Glipizide 29975-16-4, Estazolam 34911-55-2, Bupropion 36505-84-7, Buspirone 39860-99-6, Pipothiazine 42399-41-7, Diltiazem 28981-97-7, Alprazolam 30516-87-1, Zidovudine 36894-69-6, Labetalol 51481-61-9, Cimetidine 53179-11-6, Loperamide **53994-73-3**, 54910-89-3, Fluoxetine **55268-75-2**, Cefuroxime 3, Acyclovir 62571-86-2, Captopril 63590-64-7, Terazosin Cefaclor 59277-89-3, Acyclovir 66357-35-5, Ranitidine 72509-76-3, Felodipine 73590-58-6, Omeprazole 75847-73-3, Enalapril 76584-70-8, Divalproex 75330-75-5, Lovastatin sodium 76824-35-6, Famotidine 76963-41-2, Nizatidine 79617-9 Sertraline 79794-75-5, Loratidine 79902-63-9, Simvastatin 81093-37-0, Pravastatin 81103-11-9, Clarithromycin 82419-36-1, 79617-96-2, Ofloxacin 82626-48-0, Zolpidem 83799-24-0, Fexofenadine 83905-01-5, Azithromycin 85441-61-8, Quinapril 85721-33-1, Ciprofloxacin 86541-75-5, Benazepril 87333-19-5, Ramipril 88150-42-9, Amlodipine 92665-29-7, Cefprozil 97322-87-7, Troglitazone 98048-97-6, Fosinopril 110268-21-8, Opadry clear RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers) 9005-25-8, Starch, biological studies 74811-65-7, Sodium croscarmellose 9063-38-1, Sodium starch glycolate RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (diffusion enhancer; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers)

IT

IT

71010-52-1, Gellan gum

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hydrophilic carrier or diffusion enhancer; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers)

57-55-6D, Propylene glycol, esters 9000-01-5, Gum acacia IT 9000-07-1D, Carrageenan, salts or derivs. 9000-30-0, Guar gum 9000-40-2, Locust bean gum 9000-65-1, Gum tragacanth 9003-01-4, Carboxypolymethylene 9003-39-8, PVP 9004-32-4, Sodium carboxymethyl cellulose 9004-34-6D, Cellulose, derivs. 9004-42-6, Carboxyethyl cellulose 9004-62-0, Hydroxyethyl cellulose 9004-64-2, Hydroxypropyl cellulose 9004-65-3, Hydroxypropyl methylcellulose 9005-32-7D, Alginic acid, derivs. or salts 9032-42-2, Hydroxyethyl methyl cellulose 11078-30-1,

Galactomannan 11078-31-2, Glucomannan 11138-66-2, Xanthan gum 25322-68-3, Polyethylene glycol RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(hydrophilic carrier; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers)

57-11-4, Stearic acid, biological studies 57-11-4D, Stearic acid, derivs. and salts 4070-80-8, Sodium stearyl fumarate 7631-86-9, Silica, biological studies 14807-96-6, Talc, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (lubricant; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers)

77-93-0, Triethyl citrate

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (plasticizer; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers)

RE. CNT THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

(1) Bengtsson; US 5690960 A 1997 HCAPLUS

15686-71-2, Cephalexin 53994-73-3, Cefaclor

**55268-75-2**, Cefuroxime

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers)

15686-71-2 HCAPLUS RN

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-methyl-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 53994-73-3 HCAPLUS

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, (6R, 7R)- (9CI) (CA) INDEX NAME)

Absolute stereochemistry.

55268-75-2 HCAPLUS

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[[(aminocarbony])oxy]methyl]-7-[[(2Z)-2-furanyl(methoxyimino)acetyl]amin o]-8-oxo-, (6R, 7R) - (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

11078-30-1, Galactomannan IT

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hydrophilic carrier; controlled and sustained release dosage forms containing hydrophilic carriers and diffusion enhancers)

RN 11078-30-1 HCAPLUS

CN D-Galacto-D-mannan (9CI) (CA INDEX NAME)

#### \*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

ANSWER 4 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN L44

2001:814019 HCAPLUS AN

135:348920 DN

ED Entered STN: 09 Nov 2001

Modified release pharmaceutical formulations containing active principles ΤI having antibiotic activity

Maggi, Lauretta; Conte, Ubaldo IN

Laboratorio Italiano Biochimico Farmaceutico Lisapharma S.P.A., Italy

Eur. Pat. Appl., 14 pp. S<sub>0</sub>

CODEN: EPXXDW

DT Patent

English

ICM A61K009-20 IC

ICS A61K009-28; A61K031-545

63-6 (Pharmaceuticals)

ran.	PATENT NO.				KIND DATE			APPLICATION NO.							DATE			
ΡI	EP	1151				Al	_	2001			EP 2						0010	
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
						LV,												
	ΙT	2000	MI09'	72		A1		2001	1105		IT 2	000-	MI97	2		2	0000	504
	IT	1318	495			B1		20030	0825									
PRAI	ΙT	2000	-MI9	72		Α		20000	0504									
CLASS	S																	
PATI	ENT	NO.		CLA	SS	PATE	NT I	FAMILY	Y CL	ASSI	FICA	TION	COD	ES				

EP 1151747 ICM A61K009-20

ICS A61K009-28; A61K031-545 A61K009/20H6F2; A61K009/28H6F2 **ECLA** EP 1151747

Formulations are based on substances endowed with antibiotic activity for the preparation of oral hydrophilic matrixes (modified release tablets). hydrophilic matrixes can release the carried antibiotic gradually and following a release profile which can be established by suitable in vitro tests. An essential feature of said hydrophilic matrixes consists in that they can provide a high bioavailability characterized by such extended effective plasma levels to enable only one or two daily administrations, thus simplifying dosage and correct use by the patient. Coated tablets were prepared containing cefaclor monohydrate.

ST controlled release tablet antibiotic; cephalosporin controlled

release tablet

```
Drug delivery systems
         (controlled-release; modified release pharmaceutical formulations
         containing active principles having antibiotic activity)
IT
     Antibiotics
     Dissolution rate
     Drug bioavailability
     Fluidized beds
     Plasticizers
         (modified release pharmaceutical formulations containing active principles
        having antibiotic activity)
     Polyoxyalkylenes, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (modified release pharmaceutical formulations containing active principles
        having antibiotic activity)
     Drug delivery systems
IT
         (tablets, controlled-release; modified release pharmaceutical
         formulations containing active principles having antibiotic activity)
ΙT
     53994-73-3, Cefaclor
     RL: BPR (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
         (modified release pharmaceutical formulations containing active principles
        having antibiotic activity)
IT
                          7585-39-9, β-Cyclodextrin
                                                        8050-81-5,
     1398-61-4, Chitin
                    9000-07-1, Carrageenan
                                                                    9002-18-0, Agar
                                              9000-69-5, Pectin
     Simethicone
     9002-89-5 9003-39-8, Pvp 9004-32-4, Carboxymethyl cellulose 9004-53-9, Dextrin 9004-62-0, Hydroxyethyl cellulose 9004-6
                                                                   9004-65-3, Hpmc
     9004-67-5, Methyl cellulose 9012-72-0, Glucan 9012-76-4, Chitosan 9036-88-8, Mannan 9057-02-7, Pullulan 11078-30-1,
     Galactomannan 11138-66-2, Xanthan 25322-68-3, Peg 39464-87-4, Scleroglucan 106392-12-5, Poloxamer
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (modified release pharmaceutical formulations containing active principles
        having antibiotic activity)
     50-59-9, Cephaloridine 153-61-7, Cephalothin 3577-01-3,
     Cephaloglycine 10206-21-0, Cefacetrile 11111-12-9, Cephalosporin 15686-71-2, Cephalexin 21593-23-7
       Cephapirin 25953-19-9, Cephazolin 34444-01-4,
     Cefamandol 38821-53-3, Cephradine 50370-12-2
     51627-14-6, Cefatrizine 51762-05-1, Cefroxadine 55268-75-2,
     Cefuroxime 56796-20-4, Cefmetazole 60925-61-3, Ceforanide
     61270-58-4, Cefonicid 62893-19-0, Cefoperazone
                                                           63527-52-6,
                   64544-07-6, Cefuroxime axetil 64952-97-2, Moxalactam
     Cephotaxime
     70797-11-4, Cefpiramide 82219-78-1, Cefuzonam
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (modified release pharmaceutical formulations containing active principles
        having antibiotic activity)
     77-93-0, Triethyl citrate
IT
                                   77-94-1, Tributyl citrate
     Diethylphthalate 102-76-1, Triacetin 109-43-3, Dibutyl sebacate
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (plasticizer; modified release pharmaceutical formulations containing
        active principles having antibiotic activity)
RE. CNT
               THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
(1) Kim, H; WO 9949868 A 1999 HCAPLUS
(2) LI, X; HCAPLUS
(3) LI, X; GUANGDONG YAOXUEYUAN XUEBAO 2000, V16(1), P14 HCAPLUS
(4) Oren, P; US 4968508 A 1990 HCAPLUS
(5) Ranbaxy Lab Ltd; EP 0923934 A 1999 HCAPLUS
(6) Uemura, T; US 4695467 A 1987 HCAPLUS
IT 53994-73-3, Cefaclor
     RL: BPR (Biological process); BSU (Biological study, unclassified); PRP
     (Properties); THU (Therapeutic use); BIOL (Biological study); PROC
     (Process); USES (Uses)
         (modified release pharmaceutical formulations containing active principles
        having antibiotic activity)
     53994-73-3 HCAPLUS
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Page 26

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, (6R, 7R)- (9CI) (CA)INDEX NAME)

Absolute stereochemistry.

IT 11078-30-1, Galactomannan

RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL

(Biological study); USES (Uses)

(modified release pharmaceutical formulations containing active principles

having antibiotic activity)

RN 11078-30-1 HCAPLUS

D-Galacto-D-mannan (9CI) (CA INDEX NAME) CN

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

153-61-7, Cephalothin 11111-12-9, Cephalosporin 15686-71-2, Cephalexin 21593-23-7, Cephapirin

25953-19-9, Cephazolin 34444-01-4, Cefamandol

38821-53-3, Cephradine 50370-12-2 55268-75-2,

Cefuroxime 56796-20-4, Cefmetazole 62893-19-0,

Cefoperazone

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(modified release pharmaceutical formulations containing active principles

having antibiotic activity)

153-61-7 HCAPLUS

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R, 7R)- (9CI)

(CA INDEX NAME)

Absolute stereochemistry.

RN 11111-12-9 HCAPLUS

Cephalosporin (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

RN 15686-71-2 HCAPLUS

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 7-[[(2R)-aminophenylacetyl]amino]-3-methyl-8-oxo-, (6R, 7R)- (9CI)INDEX NAME)

Absolute stereochemistry.

RN 21593-23-7 HCAPLUS

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-, (6R, 7R)(9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 25953-19-9 HCAPLUS

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[[(5-methyl-1,3,4-thiadiazol-2-yl)thio]methyl]-8-oxo-7-[(1H-tetrazol-1-ylacetyl)amino]-, (6R,7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 34444-01-4 HCAPLUS

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-hydroxyphenylacetyl]amino]-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 38821-53-3 HCAPLUS

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-amino-1,4-cyclohexadien-1-ylacetyl]amino]-3-methyl-8-oxo-, (6R,7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 50370-12-2 HCAPLUS
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, (6R, 7R)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

$$\begin{array}{c|c} & & & \\ &$$

RN 55268-75-2 HCAPLUS
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-[[(2Z)-2-furanyl(methoxyimino)acetyl]amin
o]-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

RN 56796-20-4 HCAPLUS
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(cyanomethyl)thio]acetyl]amino]-7-methoxy-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, (6R, 7S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 62893-19-0 HCAPLUS
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-[[(4-ethyl-2,3-dioxo-1-piperazinyl)carbonyl]amino](4-hydroxyphenyl)acetyl]amino]-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, (6R,7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L44 ANSWER 5 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN

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1999:640708 HCAPLUS
AN
DN
      131:262650
      Entered STN: 08 Oct 1999
ED
ΤI
      Sustained-release compositions containing cefaclor
      Kim, Hyun Soo; Park, Young Joon
IN
     Yuhan Corp., S. Korea
PCT Int. Appl., 24 pp.
PA
S<sub>0</sub>
      CODEN: PIXXD2
DT
      Patent
LA
      English
      ICM A61K031-545
      ICS A61K009-22; A61K009-46
CC
      63-6 (Pharmaceuticals)
FAN. CNT 1
      PATENT NO.
                                      DATE
                                                     APPLICATION NO.
                                                                                 DATE
                              KIND
PΙ
      WO 9949868
                               A1
                                       19991007
                                                     WO 1999-KR159
                                                                                 19990401
           W: AE,
                    AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA,
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               DE, DK, EE,
                              ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
                             KP, KR, KZ, LC,
NO, NZ, PL, PT,
                                                 LK, LR, LS, LT,
                                                                    LU,
                                                                         LV,
SI,
                                                                              MD, MG, MK,
               JP, KE, KG,
                                                 RO, RU, SD,
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                         MX,
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                                                                              SK,
                                                                                   SL,
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                                                 VN,
                                                     YU, ZA, ZW,
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                        TJ,
                              TM
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                                                      UG,
          RW: GH, GM,
                              LS.
                                  MW,
                                  GR,
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      AU 9930572
                                       19991018
                                                     AU 1999-30572
                                                                                 19990401
                               A1
                                                     EP 1999-912140
      EP 1067937
                                       20010117
                                                                                 19990401
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               AT, BE, CH, DE,
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                                   FI, RO
      IP 2002509887
                               T2
                                       20020402
                                                     JP 2000-540831
                                                                                 19990401
PRAI KR 1998-11397
                                       19980401
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      WO 1999-KR159
                                       19990401
CLASS
 PATENT NO.
                    CLASS
                            PATENT FAMILY CLASSIFICATION CODES
                    ICM
 WO 9949868
                             A61K031-545
     ICS A61K009-22; A61K009-46
9949868 ECLA A61K009/00L2; A61K033/00
A composition comprising 30-90 % cefaclor, 5-60 % a hydroswelling polymer, and 1-10 % a salt based on the total weight of the composition releases cefaclor in
 WO 9949868
      the gastric fluid in a steady sustained manner for > 20 h. Cefaclor 400 g
      was mixed with hydroxypropyl Me cellulose 20, CaCO3 20, NaHCO3 10, citric
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acid 10, hydroxypropyl Me cellulose phthalate 10, and colloidal silica 2 g. The mixture was granulated and the obtained granules were mixed with hydroxypropyl Me cellulose 60, CaCO3 20, NaHCO3 5, Na CMC 30, colloidal silica 1, and Mg stearate 3 g. The resulting mixture was formed into tablets. When the tablet thus obtained was added to an artificial fluid, it floated to the upper layer and remained afloat for > 6 h. The cumulative dissoln. rate of cefaclor from the tablet in 0.1 N HCl aqueous

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solution at 37^\circ was 18.03, 25.58, 37.17, 59.52, 91.24, and 100 % at 15, 30, 60, 120, 240, and 480 min, resp.
      sustained release tablet cefaclor cellulose ether
      Dissolution rate
           (sustained-release oral composition containing cefaclor and water-swellable
          polymers and (bi) carbonates and acids)
IT
      Drug delivery systems
           (tablets, sustained-release; sustained-release oral composition containing
      cefaclor and water-swellable polymers and (bi) carbonates and acids) 77-92-9, biological studies 87-69-4, biological studies 110-16-7, 2-Butenedioic acid (2Z)-, biological studies 144-55-8, Sodium
      bicarbonate, biological studies 298-14-6, Potassium bicarbonate
      471-34-1, Calcium carbonate, biological studies 497-19-8, Sodium
      carbonate, biological studies 533-96-0, Sodium sesquicarbonate 546-93-0, Magnesium carbonate 584-08-7, Potassium carbonate 9003-39-8, PVP 9004-32-4, Sodium CMC 9004-64-2, Hydroxypropyl
      cellulose 9004-65-3, Hydroxy alginate 53994-73-3, Cefaclor
                     9004-65-3, Hydroxypropyl methyl cellulose 9005-38-3, Sodium
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
           (sustained-release oral composition containing cefaclor and water-
          swellable polymers and (bi) carbonates and acids)
RE. CNT
                  THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
    Eisai Co Ltd; EP 0235718 A2 1987 HCAPLUS
    Eli Lilly And Company; EP 0280571 A2 1988 HCAPLUS
Shionogi Seiyaku Kabushiki Kaisha; GB 2141342 A 1984 HCAPLUS
      9003-39-8, PVP 53994-73-3, Cefaclor
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
          (sustained-release oral composition containing cefaclor and water-swellable polymers and (bi) carbonates and acids)
      9003-39-8 HCAPLUS
CN
      2-Pyrrolidinone, 1-ethenyl-, homopolymer (9CI) (CA INDEX NAME)
      CM
      CRN 88-12-0
      CMF C6 H9 N O
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RN 53994-73-3 HCAPLUS CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, (6R,7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L44 ANSWER 6 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN AN 1999:42584 HCAPLUS

DN 130:105323

ED Entered STN: 21 Jan 1999

TI Control of acidic gut syndrome with an agent controlling acid and endotoxin accumulation in the gastrointestinal tract

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Rowe, James Baber
PA
     Australia
S0
     PCT Int. Appl., 59 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
     ICM A61K031-71
IC
     ICS
         A61K038-43; A61K038-46; A61K038-47
CC
     1-9 (Pharmacology)
     Section cross-reference(s): 63
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     PATENT NO.
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                                                                      DATE
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                                                                      19980626
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                                 19990107
     CA 2294904
                           AA
                                              CA 1998-2294904
                                                                      19980626
     AU 9880931
                                 19990119
                                              AU 1998-80931
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                           A1
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     EP 1017402
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     JP 2002511865
                                              JP 1999-505159
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     NZ 502445
                                 20030328
                                              NZ 1998-502445
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                                              US 2000-446801
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     US 6303572
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                           R1
     US 6468964
                           B1
                                 20021022
                                              US 2001-912886
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PRAI AU 1997-7582
                                 19970627
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CLASS
 PATENT NO.
                 CLASS
                        PATENT FAMILY CLASSIFICATION CODES
 WO 9900136
                  ICM
                         A61K031-71
                         A61K038-43; A61K038-46; A61K038-47
                 ICS
 WO 9900136
                 ECLA
                         A61K038/12
                         A61K038/12
 US 6303572
                 ECLA
 US 6468964
                 ECLA
                         A61K038/12
    A method is provided for the treatment or prophylaxis of acidic gut
     syndrome resulting from the accumulation of acid and production of endotoxin
     in the gastrointestinal tract of a human or an animal, the accumulation
     resulting from the fermentation of carbohydrate in the gastrointestinal tract of
     the human or animal. The method comprises administering to said human or
     animal an effective amount of an active agent capable of preventing or
     controlling acid and endotoxin accumulation in the gastrointestinal tract.
ST
     acidic gut syndrome acid endotoxin accumulation modulation
IT
     Intestine, disease
        (Crohn's, acidic gut syndrome-associated; agent controlling acid and
        endotoxin accumulation in GI tract for acidic gut syndrome control)
IT
     Endocrine system
     Kidney
     Pancreas
     Thyroid gland
        (acidic gut syndrome-associated effect on; agent controlling acid and
        endotoxin accumulation in GI tract for acidic gut syndrome control)
     Alzheimer's disease
IT
     Arthritis
     Asthma
     Cachexia
     Cystic fibrosis
     Dermatitis
     Diabetes mellitus
     Hypertension
     Mastitis
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Multiple sclerosis
     Myasthenia gravis
     Neoplasm
     Osteoarthritis
     Osteoporosis
     Rheumatoid arthritis
        (acidic gut syndrome-associated; agent controlling acid and endotoxin
        accumulation in GI tract for acidic gut syndrome control)
IT
     Acidosis
     Alfalfa (Medicago sativa)
     Anaerovibrio
     Antibiotics
     Bacteria (Eubacteria)
     Bacteroides
     Barley
     Butyrivibrio
     Cottonseed meal
     Drug targeting
     Gram-positive bacteria (Firmicutes)
     Immune system
     Ions
     Megasphaera
     Peptococcus
     Propionibacterium
     Selenomonas
     Sheep
     Succinivibrio
     Succinomonas
     Swine
     Vaccines
     Veillonella
     Нq
        (agent controlling acid and endotoxin accumulation in GI tract for
        acidic gut syndrome control)
    Bentonite, biological studies
     Clays, biological studies
     Enzymes, biological studies
     Smectite-group minerals
     Zeolite-group minerals
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (agent controlling acid and endotoxin accumulation in GI tract for
        acidic gut syndrome control)
  Acids, biological studies
     Cytokines
     Limestone, biological studies
     Lipopolysaccharides
     Polysaccharides, biological studies
     Tumor necrosis factors
    RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
        (agent controlling acid and endotoxin accumulation in GI tract for
        acidic gut syndrome control)
ΙT
    Antibiotics
        (aminoglycoside; agent controlling acid and endotoxin accumulation in
        GI tract for acidic gut syndrome control)
ΙT
    Nervous system
        (amyotrophic lateral sclerosis, acidic gut syndrome-associated; agent
        controlling acid and endotoxin accumulation in GI tract for acidic gut
        syndrome control)
     Ionophores
IT
        (antibiotic; agent controlling acid and endotoxin accumulation in GI
        tract for acidic gut syndrome control)
IT
    Glycolipids
     Glycopeptides
     Peptides, biological studies
     Polythiazoles
     Proteins, general, biological studies
     Sulfonamides
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Qazi 10/642194

Page 33

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (antibiotic; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT (appendicitis, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) ΙT (bacterial, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) ΙT Fermentation (carbohydrate; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Tooth (caries, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IΤ Fatigue, biological (chronic fatigue syndrome, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Intestine, disease IΤ (colitis, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Flours and Meals (corn, hind gut fermentable diet containing; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Respiratory tract (disease, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Digestive tract (disease, acidic gut syndrome; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IΤ (disease, infection, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Homeostasis Immunity (disorder, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Toxins RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (endotoxins; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Drug delivery systems (enteric-coated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Carbohydrates, biological studies RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (fermentation; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) (flour, hind gut fermentable diet containing; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Digestive tract (gastroenteritis, immune condition associated with, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Stress, animal (heat, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Infection (helminth or microbial, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome

control) Lung, disease

(hemorrhage, post-exercise, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) ΙT Rice (Oryza sativa) (hind gut fermentable diet containing; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Electrolytes, biological (imbalance, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Minerals, biological studies RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) ΙT (imbalance, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) ΙT Aerococcus Alloiococcus Carnobacterium Enterococcus Lactobacillus Lactococcus Leuconostoc Pediococcus Streptococcus Streptococcus bovis Tetragenococcus (immunization against; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Wool (impaired growth, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Reproductive tract (impaired reproductive performance, acidić gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Fungi ΙT Herpesviridae Parasitic worm Protozoa (infection, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT (intake reduction, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Intestine, disease IT (irritable bowel syndrome, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Antibiotics (macrolide; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Enzymes, biological studies RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (pectic; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Opioids RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (peptide; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Uronic acids RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (polyuronic acids; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Digestive tract Digestive tract

Stomach, disease (ulcer, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Intestine, disease (ulcerative colitis; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Infection (viral, acidic gut syndrome-associated; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Fatty acids, biological studies RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (volatile; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) Galactosides RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process) (α-galactosides; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) IT Lactams RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (β-, antibiotics; agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) 56-75-7D, Chloramphenicol, derivs. 57-62-5, Chlortetracycline 59-87-0, IT Nitrofurazone 61-32-5, Methicillin 66-79-5, Oxacillin 67-20-9, Nitrofurantoin 67-45-8, Furazolidone 69-53-4, Ampicillin 79-57-2, Oxytetracycline 87-08-1, Penicillin V 114-07-8, Erythromycin 154-21-2, Lincomycin 804-36-4, Nitrovin 1066-17-7, Polymyxin E 1318-00-9, Vermiculite 1318-63-4, Heulandite 1318-74-7, Kaolinite, biological studies 1318-93-0, Montmorillonite, biological studies 1319-41-1, Saponite 1393-48-2, Thiostrepton 1401-69-0, Tylosin 1404-26-8, Polymyxin B 1404-90-6, Vancomycin 1405-89-6, Bacitracin zinc 1406-05-9, Penicillin 1476-53-5, Novobiocin sodium 1695-77-8, Spectinomycin 8025-81-8, Spiramycin 9000-92-4, Amylase 9001-22-3, Emulsin 9001-42-7, \alpha-Glucosidase 9001-57-4, Invertase 9001-92-7, Protease 9002-07-7, Trypsin 9001-75-6, Pepsin Trypsinogen 9004-07-3, Chymotrypsin 9011-97-6, Cholecystokinin 9012-54-8, Cellulase 9015-78-5, Glucanase 9025-35-8, 9032-08-0, Amyloglucosidase 9040-13-5, 9074-98-0, β-Glucanase 11006-76-1, Virginiamycin α-Galactosidase Virginiamycin M 11006-76-1D, Virginiamycin, M1 fraction analogs 11015-37-5, Flavomycin 11017-43-9, Siomycin 11054-70-9, Lasalocid 11111-12-9, Cephalosporin 11115-82-5, Enramycin 12001-29-5, Chrysotile 12161-84-1, Lizardite 12172-85-9, Beidellite 12173-10-3, Clinoptilolite 12173-60-3, Illite 12174-11-7, Palygorskite 12269-78-2, Pyrophyllite 12298-43-0, Halloysite 12609-84-6, Thiopeptin 12650-69-0, Mupirocin 14807-96-6, Talc, biological studies 16846-24-5, Josamycin 18323-44-9, Clindamycin 23152-29-6, Virginiamycin S1 37244-77-2, Sporangiomycin 37278-89-0, Xylanase 37332-99-3, Avoparcin 49648-37-5, Fluoramphenicol 51667-26-6D, Oxazolidinone, derivs. 53003-10-4, Salinomycin 53024-98-9D, Everninomycin, derivs. 55134-13-9, Narasin 55297-95-5, Tiamulin 55852-84-1, Bacitracin methylene disalicylate 56377-79-8, Nosiheptide 61036-62-2, Teicoplanin 65454-16-2, Taitomycin 65454-59-3, Sulfomycin 70324-48-0, Bottromycin 65454-16-2, Taitomycin 65454-59-3, Sulfomycin 70324-48-0, Botartrate 75139-06-9, Tetronasin 79805-24-6, β-Casomorphin 82419-36-1, Ofloxacin 85721-33-1, Ciprofloxacin 100986-85-4, 105956-97-6, Clinafloxacin 109545-84-8, SCH 27899 Levofloxacin 113041-69-3, Magainin 117742-13-9, Ardacin 126602-89-9, RP 59500 127254-12-0, DU 6859 110871-86-8, Sparfloxacin 119914-60-2, Grepafloxacin 127254-12-0, DU 6859a 147059-72-1, Trovafloxacin 165800-03-3, U-100766 219686-62-1, 219686-64-3, U 100572 Thiopeptone RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (agent controlling acid and endotoxin accumulation in GI tract for acidic gut syndrome control) 50-21-5, biological studies 57-13-6, Urea, biological studies IT Galactose, biological studies 64-19-7, Acetic acid, biological studies

79-09-4, Propanoic acid, biological studies 79-33-4, biological studies 107-92-6, Butyric acid, biological studies 147-81-9, Arabinose

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9012-72-0, Glucan
     9005-25-8, Starch, biological studies
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     Arabinogalactan 9037-55-2, Galactan 9040-27-1, Arabinoxylan
     9041-22-9, β-Glucan 9046-38-2, Galacturonan 9060-75-7, L-Arabinan
     10326-41-7, D-Lactic acid, biological studies 11078-30-1,
     Galactomannan 37294-28-3, Xyloglucan 39280-21-2,
     Rhamnogalacturonan
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
         (agent controlling acid and endotoxin accumulation in GI tract for
         acidic gut syndrome control)
     27194-24-7D, Nitrofuran, derivs.
IT
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
         (antibiotic; agent controlling acid and endotoxin accumulation in GI
         tract for acidic gut syndrome control)
ΙT
     187112-48-7, Raftilose
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
         (hind gut fermentable diet containing; agent controlling acid and endotoxin
        accumulation in GI tract for acidic gut syndrome control)
     80738-43-8, Lincosamide
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
         (lincosamide antibiotics; agent controlling acid and endotoxin
        accumulation in GI tract for acidic gut syndrome control)
     9073-60-3, Penicillinase
IΤ
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
         (penicillins resistant to; agent controlling acid and endotoxin
        accumulation in GI tract for acidic gut syndrome control)
     512-69-6, Raffinose
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL
     (Biological study); PROC (Process)
         (raffinose-series galactosides; agent controlling acid and endotoxin
        accumulation in GI tract for acidic gut syndrome control)
4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE. CNT
(1)
    Muir; Journal of Animal Science 1980, V50(3), P547 HCAPLUS
(2) Nagaraja; Journal of Animal Science 1981, V53(1), P206 HCAPLUS (3) Rowe; AU 3228489 B 1989
(4) Rowe; AU 4324596 A 1996
     11111-12-9, Cephalosporin
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
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        acidic gut syndrome control)
     11111-12-9 HCAPLUS
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     Cephalosporin (9CI)
                           (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     11078-30-1, Galactomannan
ΙT
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
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11078-30-1 HCAPLUS
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L44 ANSWER 7 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN
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     130:17234
     Entered STN: 23 Nov 1998
ED
     Preparation of microsphere drug delivery systems
ΤI
     Wu, Xiao Yu; Liu, Zhi
IN
PA
     Can.
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PCT Int. Appl., 47 pp.
S0
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     ICM A61K009-16
     ICS
         A61K047-48; A61K045-06; A61K031-475; A61K031-27
     63-6 (Pharmaceuticals)
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PRAI US 1997-45710P
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     WO 1998-CA419
                                  19980506
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                         PATENT FAMILY CLASSIFICATION CODES
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                         A61K047-48; A61K045-06; A61K031-475; A61K031-27
A61K009/16H6B; A61K009/16H6F; A61K009/16K;
A61K031/475+M; A61K045/06; A61K047/48W8
                  ICS
 WO 9850018
                  ECLA
     A drug delivery composition comprising microspheres containing at least one
     chemotherapeutic agent and at least 1 chemosensitizer wherein the
     microspheres have a biodegradable polymer matrix with functional groups
     which associate with the chemotherapeutic agent and chemosensitizer is
     described. Carboxymethyl dextran microspheres were prepared and mixed with
     1% verapamil or doxorubicin aqueous solution The microspheres showed sustained
     drug release.
     microsphere drug delivery system polymer
     Calmodulins
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (antagonists; preparation of polymeric microsphere drug delivery systems)
     Polymers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (biodegradable; preparation of polymeric microsphere drug delivery systems)
     Ion channel blockers
IT
         (calcium; preparation of polymeric microsphere drug delivery systems)
IT
     Polyesters, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (caprolactone-based; preparation of polymeric microsphere drug delivery
        systems)
IT
     Polyesters, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (hydroxycarboxylic acid-based; preparation of polymeric microsphere drug
        delivery systems)
     Alkaloids, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (indole; preparation of polymeric microsphere drug delivery systems)
ΙT
     Polyesters, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (lactic acid-based; preparation of polymeric microsphere drug delivery
        systems)
     Drug delivery systems
IT
         (microspheres, sustained-release; preparation of polymeric microsphere drug
        delivery systems)
ΙT
     Polyethers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (ortho ester group-containing; preparation of polymeric microsphere drug
        delivery systems)
IT
     Antibiotics
     Antitumor agents
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Ion exchangers
          (preparation of polymeric microsphere drug delivery systems)
IT
      Albumins, biological studies
      Alkaloids, biological studies
      Glycosaminoglycans, biological studies
      Hormones, animal, biological studies
      Peptidoglycans
      Polyamides, biological studies
      Polyanhydrides
      Polyesters, biological studies
      Polysaccharides, biological studies
      Polyurethanes, biological studies
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
          (preparation of polymeric microsphere drug delivery systems)
IT
      216099-82-0P
      RL: PRP (Properties); SPN (Synthetic preparation); THU (Therapeutic use);
      BIOL (Biological study); PREP (Preparation); USES (Uses)
          (preparation of polymeric microsphere drug delivery systems)
IΤ
      RL: RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use);
      BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
          (preparation of polymeric microsphere drug delivery systems)
IT
      9005-80-5, Inulin
      RL: RCT (Reactant); THU (Therapeutic use); BIOL (Biological study); RACT
      (Reactant or reagent); USES (Uses)
          (preparation of polymeric microsphere drug delivery systems)
ΙT
      9044-05-7P, Carboxymethyl dextran
      RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological
      study); PREP (Preparation); USES (Uses)
          (preparation of polymeric microsphere drug delivery systems)
      50-07-7, Mitomycin C 50-53-3, Chlorpromazine, biological studies 52-53-9, Verapamil 56-54-2, Quinidine 57-22-7, Vincristine 85
IT
                                                                                       83-89-6,
      Quinacrine 117-89-5, Trifluoperazine
                                                         130-95-0, Quinine 865-21-4,
                      1398-61-4, Chitin 9000-07-1D, Carrageenan, analogs
      Vinblastine
      9000-65-1, Tragacanth gum 9000-69-5D, Pectinic acid, analogs 9002-18-0, Agar 9004-34-6, Cellulose, biological studies 90
                                                                                  9004-54-0,
      Dextran, biological studies 9004-61-9, Hyaluronic acid 9005-25-8, Starch, biological studies 9005-32-7, Alginic acid 9005-32-7D, Alginic
      acid, analogs 9005-49-6, Heparin, biological studies 9005-79-2, Glycogen, biological studies 9007-28-7D, Chondroitin sulfate, analogs
      9012-36-6, Agarose 9012-72-0D, Glucan, analogs 9014-63-5D, Xylan,
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      analogs
      analogs
      9040-29-3D, GalactoGlucomannan, analogs 9041-38-7D, Teichoic
                         9046-38-2D, Galacturonan, analogs
                                                                      9046-40-6D, Pectic
      acid, analogs
                          9056-36-4, Keratan sulfate
                                                              10540-29-1, Tamoxifen
      acid, analogs
      11078-27-6D, Arabinan, analogs 11078-30-1D,
      Galactomannan, analogs 11078-31-2D, Glucomannan, analogs
      11138-66-2, Xanthan gum 15802-18-3D, 2-Cyanoacrylic acid, alkyl esters, polymers 20830-81-3, Daunorubicin 21829-25-4, Nifedipine 23214-92-8, Doxorubicin 24937-72-2, Poly(maleic anhydride) 24967-94-0, Dermatan
      sulfate 24980-41-4, Poly(&-caprolactone) 25086-15-1,
      Methacrylic acid-methyl methacrylate copolymer 25248-42-4,
      Poly[oxy(1-oxo-1,6-hexanediy1)] 26009-03-0, Poly(Glycolic acid) 26023-30-3, Poly[oxy(1-methyl-2-oxo-1,2-ethanediy1)] 26100-51-6
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      Poly (lactic acid) 26124-68-5, Poly (Glycolic acid) 26700-71-0D, Poly (L-glutamine), hydroxyalkyl derivs. 28728-97-4, Poly (hydroxybutyric acid), SRU 29323-51-1 34346-01-5, Glycolic acid-lactic acid copolymer 37294-28-3D, Xyloglucan, analogs 42399-41-7, Dilitic 2020 10.
      Poly (hydroxybutyric acid) 55985-32-5, Nicardipine 62893-19-0, Cefoperazone 72509-76-3, Felodipine 73384-59-5, Ceftriaxone
      80137-67-3, &-Caprolactone-lactic acid copolymer
                                                                      128171-16-4,
      Hydroxybutyric acid-hydroxyvaleric acid copolymer
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      176442-72-1
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
          (preparation of polymeric microsphere drug delivery systems)
                 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE. CNT
(1) Behringwerke Ag; EP 0595133 A 1994 HCAPLUS
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(2) Biovecteurs As; FR 2702160 A 1994 HCAPLUS
   John, K; WO 9709036 A 1997 HCAPLUS
    Patrinove; CH 681780 A 1993 HCAPLUS
(5) Rene, F; US 4666641 A 1987 HCAPLUS
(6) Rolf, K; US 5057304 A 1991 HCAPLUS
(7) Univ Texas; WO 9402106 A 1994 HCAPLUS
     9040-29-3D, GalactoGlucomannan, analogs 11078-30-1D, Galactomannan, analogs 62893-19-0, Cefoperazone
     73384-59-5, Ceftriaxone
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         (preparation of polymeric microsphere drug delivery systems)
     9040-29-3 HCAPLUS
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     D-Galacto-D-gluco-D-mannan (9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     11078-30-1 HCAPLUS
CN
     D-Galacto-D-mannan (9CI) (CA INDEX NAME)
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     62893-19-0 HCAPLUS
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
CN
     7-[[(2R)-[[(4-ethyl-2, 3-dioxo-1-piperazinyl)carbonyl]amino](4-hydroxyphenyl)acetyl]amino]-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-
     oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)
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Absolute stereochemistry.

RN 73384-59-5 HCAPLUS CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-3-[[(1,2,5,6-tetrahydro-2-methyl-5,6-dioxo-1,2,4-triazin-3-yl)thio]methyl]-, (6R,7R)- (9Cl) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

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L44 ANSWER 8 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN
     1998:580026 HCAPLUS
AN
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     129:207213
     Entered STN: 11 Sep 1998
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ΤI
     Combination agent for oral administration of antibiotics
IN
     Posanski, Ulrich
PA
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     Ger. Offen., 4 pp.
S0
     CODEN: GWXXBX
DT
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LA
     German
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          A61K031-70; A61K031-495
     A61K031-43, A61K031-70, A61K031-495
ICI
CC
     63-6 (Pharmaceuticals)
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                                                APPLICATION NO.
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CLASS
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 WO 9836732
                  ECLA
                          A61K009/00N2; A61K031/43+M; A61K031/70R5L
     An aqueous suspension for oral administration of antibiotics comprises (a)
     antibiotic-containing particles with a polymeric, cationic, permeable,
     swellable and/or gastric juice-soluble coating and (b) a syrup base with pH
     5-10; the 2 components are packaged sep. and combined immediately before
     administering the 1st dose to improve the stability of the composition during
     storage. The particle coating masks the bitter flavor of the antibiotic
     for pediatric use. Thus, a mixture of amoxicillin-3H2O 62.5, K clavulanate
     32.813, and Avicel PH 101 15.0 kg was spray-coated with 150 kg 15 weight% Eudragit E solution in iso-PrOH-Me2CO (80:20) in an N2 atmospheric, sieved, dried, mixed with 5.5 kg Syloid 244 FP, and dispensed in 11.065-g portions into
     sachets. A syrup base was prepared by dissolving Na saccharin 3, NaH2PO4
     50, and Na2HPO4 950 g in 70 kg Karion FP at 40^{\circ} , adjusting the pH
     to 7.2, adding 5 g raspberry flavoring, and dispensing into screw-capped
     brown glass bottles.
     antibiotic coated particle oral syrup
     Antibiotics
         (combination agent for oral administration of antibiotics)
IT
     Polymers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (films, permeable, swellable; combination agent for oral
        administration of antibiotics)
     Drug delivery systems
ΙT
         (particles, coated; combination agent for oral administration of
        antibiotics)
IT
     Films
         (polymeric, permeable, swellable; combination agent for oral
        administration of antibiotics)
IT.
     Drug delivery systems
```

(syrups; combination agent for oral administration of antibiotics) 79-10-7D, 2-Propenoic acid, polymers, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) IT (coatings; combination agent for oral administration of antibiotics) 69-53-4, Ampicillin 132-98-9, Phenoxymethylpenicillin potassium IT 985-16-0, Sodium nafcillin 1245-44-9, Propicillin potassium 1405-10-3, Neomycin sulfate 3116-76-5, Dicloxacillin 15686-71-2, Cephalexin 26787-78-0, Amoxicillin 28657-80-9, C 50370-12-2, Cefadroxil 50972-17-3, Bacampicillin 28657-80-9, Cinoxacin 51940-44-4, Pipemidic acid 53994-73-3, Cefaclor 55268-75-2, 61177-45-5, Potassium clavulanate 61336-70-7, Amoxicillin 70458-96-7, Norfloxacin 74011-58-8, Enoxacin 76470-66-1, Cefuroxime trihydrate Loracarbef 79198-29-1, Amoxicillin-clavulanic acid mixture 79350-37-1, Cefixime 79660-72-3, Fleroxacin 80210-62-4, 81103-11-9, Clarithromycin 82419-36-1, Ofloxacin Cefpodoxime 83905-01-5, Azithromycin 85721-33-1, Ciprofloxacin 87239-81-4, Cefpodoxime-proxetil 92665-29-7, Cefprozil 98079-51-7, Lomefloxacin 147059-72-1, Trovafloxacin RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (combination agent for oral administration of antibiotics)

79-10-7D, 2-Propenoic acid, alkyl esters, polymers with methacrylic acid (esters), biological studies 79-41-4D, alkyl esters, polymers with acrylic acid (esters) 9003-21-8, Poly(methyl acrylate) 9003-32-1, Poly(ethyl acrylate) 9011-87-4, Methyl acrylate/methyl methacrylate 24938-16-7, Eudragit E 25751-21-7, Acrylic acid/methacrylic copolymer acid copolymer 56399-02-1, Ethyl acrylate/ethyl methacrylate copolymer 107500-57-2 211863-54-6 212058-93-0, Karion FP RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (combination agent for oral administration of antibiotics)

15686-71-2, Cephalexin 50370-12-2, Cefadroxil IT

53994-73-3, Cefaclor 55268-75-2, Cefuroxime 79350-37-1, Cefixime

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES

(combination agent for oral administration of antibiotics)

RN15686-71-2 HCAPLUS

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-methyl-8-oxo-, (6R, 7R)- (9CI)INDEX NAME)

Absolute stereochemistry.

50370-12-2 HCAPLUS RN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, (6R, 7R)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 53994-73-3 HCAPLUS

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 55268-75-2 HCAPLUS

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-[[(2Z)-2-furanyl(methoxyimino)acetyl]amin
o]-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

RN 79350-37-1 HCAPLUS

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2Z)-(2-amino-4-thiazolyl)[(carboxymethoxy)imino]acetyl]amino]-3-ethenyl-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.

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ANSWER 9 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN
L44
     1998:123996 HCAPLUS
AN
DN
     128:184696
     Entered STN: 28 Feb 1998
ED
TI
     Easy to swallow oral medicament composition
     Gruber, Peter
IN
PA
     Losan Pharma G.m.b.H., Germany; Gruber, Peter
     PCT Int. Appl., 65 pp. CODEN: PIXXD2
S<sub>0</sub>
DT
     Patent
LA
     German
     ICM A61K009-50
IC
     ICS A61K009-54
CC
     63-6 (Pharmaceuticals)
FAN. CNT 1 3
     PATENT NO.
                                   DATE
                                                APPLICATION NO.
                                                                         DATE
                           KIND
                                                                         19970814
PΙ
     WO 9806385
                            A1
                                   19980219
                                                WO 1997-CH299
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                           CA, CN,
                                                                       SK, TR, UA,
                                                                       MC,
                                                                           NL, PT, SE
          RW: AT, BE, CH, DE, DK,
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                                   19980219
                                                CA 1997-2262595
                                                                         19970814
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                                   19980306
                                                AU 1997-36912
                            A1
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                                   19990602
                                                EP 1997-933611
                                                                         19970814
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     EP 918513
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                            B1
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              IE, FI
     JP 2000516222
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                                   20001205
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                                                                         19970814
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                                                AT 1997-933611
                                                                         19970814
                            E
     US 2002068088
                                   20020606
                                                US 1999-242167
                                                                         19990210
                            A 1
     US 6709678
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                                                                         20031112
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PRAI CH 1996-2006
                                   19960815
                            Α
     WO 1997-CH299
                                   19970814
     US 1999-242167
                            A1
                                   19990210
CLASS
                          PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                  CLASS
 WO 9806385
                   ICM
                          A61K009-50
                   ICS
                          A61K009-54
                          A61K009/00M18B; A61K009/00N2; A61K009/50K
A61K009/00M18B; A61K009/00N2; A61K009/50K
A61K009/00M18B; A61K009/00N2; A61K009/50K
 WO 9806385
                  ECLA
 US 2002068088
                  ECLA
 US 2004247675
                  ECLA
     An easy-to-swallow pharmaceutical composition consists of ≥1 coated
     particles with a core which contains an active substance and a coat with
     ≥1 layers. The coating layer(s) contains ≥1 hydratable,
     pharmaceutically acceptable polymer which, on contact with saliva or
     water, forms a coherent, moldable, viscous mass with a slippery surface
     which does not adhere to the mucous membranes of the mouth, and which
     prevents the active substance-containing particles from leaving the mass and
     releasing the active substance in the mouth cavity. The (outermost)
     coating layer contains ≥1 salivation-promoting agent. The
     properties of the coating make the composition suitable for administering
     highly dosed or bad-tasting active substances and even for swallowing
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without any liquid Thus, a solution of ciprofloxacin 2000, Crospovidone XL-M
      110, PVP K90 60, water 900, and EtOH 1800 g was spray-coated onto sucrose
      crystals 0.3-0.6 mm in diameter to produce core particles, which were then
      coated first with a powdered mixture of NaCl 50, Na saccharin 50, and Na
      carboxymethylstarch 50 g, and finally [after moistening with EtOH-H20
      (1:1)] with a powdered mixture of Na CM-cellulose 275 and talc 75 g.
ST
      oral pharmaceutical swallowing polymer coating; polysaccharide coating
      oral pharmaceutical swallowing
     Gelatins, biological studies
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(coatings, hydration of; easy-to-swallow oral medicament composition)
IT
      Polymers, biological studies
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
          (coatings; easy-to-swallow oral medicament composition)
IT
      Osmolytes
          (easy-to-swallow oral medicament composition)
ΙT
      Carboxylic acids, biological studies
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
          (easy-to-swallow oral medicament composition)
      Hydration, chemical
IT
          (of polymer coating; easy-to-swallow oral medicament composition)
IT
      Drug delivery systems
      Drug delivery systems
          (oral, controlled-release; easy-to-swallow oral medicament composition)
IT
      Drug delivery systems
      Drug delivery systems
          (pellets, controlled-release; easy-to-swallow oral medicament composition)
IT
      Saliva
          (polymer coating hydration by; easy-to-swallow oral medicament composition)
IT
      Coating materials
          (polymers; easy-to-swallow oral medicament composition)
IT
      Drug delivery systems
          (tablets, controlled-release; easy-to-swallow oral medicament composition)
      9000-01-5, Gum arabic
IΤ
                                    9000-30-0, Guar gum 9000-69-5, Pectin
                                           9003-39-8, PVP
      9003-01-4, Polyacrylic acid
                                                                9004-32-4, Sodium
     CM-cellulose 9004-62-0, Hydroxyethylcellulose 9004-64-2, Hydroxypropylcellulose 9004-65-3, Hydroxypropylmethylcellulose
      9004-67-5, Methylcellulose 9005-32-7, Alginic acid 11078-30-1, Galactomannan 11138-66-2, Xanthan 39421-75-5, Hydroxypropyl
      guar gum
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
          (coating, hydration of; easy-to-swallow oral medicament composition)
     79-10-7D, Acrylic acid, esters, polymers 9005-32-7D, Alginic acid, salts RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (coatings, hydration of; easy-to-swallow oral medicament composition)
IT
      50-48-6, Amitriptyline 50-78-2, Acetylsalicylic acid 52-53-9,
IT
                   57-27-2, Morphine, biological studies 58-55-9, Theophylline, studies 60-54-8, Tetracycline 89-57-6, Mesalazine
      Verapamil
      biological studies
      103-90-2, Paracetamol
                                   114-07-8, Erythromycin 146-22-5, Nitrazepam
      603-50-9, Bisacodyl 616-91-1, N-Acetyl-L-cysteine 4618-18-2, Lactulose
     8049-47-6, Pancreatin 15307-86-5, Diclofenac 15687-27-1, Ibuprofen 15722-48-2, Olsalazine 16051-77-7, Isosorbide mononitrate 18683-91-Ambroxol 21829-25-4, Nifedipine 22204-53-1, Naproxen 26787-78-0, Amoxicillin 27203-92-5, Tramadol 29122-68-7, Atenolol 36322-90-4,
                                                                                    18683-91-5.
      Piroxicam
                     42399-41-7, Diltiazem 50370-12-2, Cefadroxil
      50679-08-8, Terfenadine
                                      51333-22-3, Budesonide
                                                                      51481-61-9, Cimetidine
      53179-11-6, Loperamide 53994-73-3, Cefaclor 54182-58-0,
                                                     68844-77-9, Astemizole
                      66357-35-5, Ranitidine
      Sucralfate
                                                                                     70458-96-7
                       73590-58-6, Omeprazole 76824-35-6, Famotidine izatidine 82419-36-1, Ofloxacin 85721-33-1, Ciprofloxacin
      Norfloxacin
      76963-41-2, Nizatidine
     93107-08-5, Ciprofloxacin hydrochloride 102625-70-7, Pantoprazole RL: BAC (Biological activity or effector, except adverse); BSU (Biological
      study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
      (Uses)
          (easy-to-swallow oral medicament composition)
      50-70-4, Sorbitol, biological studies 50-81-7, L-Ascorbic acid,
      biological studies 50-99-7, D-Glucose, biological studies 57-48-7,
     D-Fructose, biological studies 57-50-1, Sucrose, biological studies 69-65-8, D-Mannitol 77-92-9, Citric acid, biological studies 87-69-4, Tartaric acid, biological studies 87-99-0, Xylitol 124-04-9, Adipic
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acid, biological studies 134-03-2, Sodium ascorbate 585-09-1, Potassium malate 585-88-6, Maltitol 676-46-0, Sodium malate 994-36-5 6915-15-7, Malic acid 7647-14-5, Sodium chloride, biological studies 9004-34-6D, Cellulose, ethers, biological studies 9063-38-1, Sodium carboxymethylstarch 14475-11-7, biological studies 15421-15-5, Potassium ascorbate 18996-35-5, Monosodium citrate 40968-90-9, Potassium tartrate, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses). (easy-to-swallow oral medicament composition) 9004-34-6, Cellulose, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) IT (microcryst.; easy-to-swallow oral medicament composition) RE. CNT THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD RE

- Benzon Pharma AS; US 5288500 A 1988 HCAPLUS (1)
- (2) Benzon Pharma AS; WO 8806893 A 1988 HCAPLUS

(3) Ko; WO 9301800 A 1993 HCAPLUS

Rhodia Farma Ltda; BR 9403617 A 1996 HCAPLUS

- (5) Showa Yakuhin Kako Co Ltd; EP 0662320 A 1995 HCAPLUS
- (6) Zyma Sa; EP 0281513 A 1988 HCAPLUS
- (7) Zyma Sa; US 4882169 A 1988 HCAPLUS

11078-30-1, Galactomannan

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (coating, hydration of; easy-to-swallow oral medicament composition) 11078-30-1 HCAPLUS

RN

D-Galacto-D-mannan (9CI) (CA INDEX NAME) CN

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

50370-12-2, Cefadroxil 53994-73-3, Cefaclor RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(easy-to-swallow oral medicament composition)

50370-12-2 HCAPLUS RN

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, (6R, 7R)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 53994-73-3 HCAPLUS

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

L44 ANSWER 10 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN AN 1996:544057 HCAPLUS

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DN
     125:185914
ED
     Entered STN: 12 Sep 1996
     Prevention of adverse behavior, diarrhea, skin disorders and infections of
     the hind gut associated with acidic conditions in humans and animals
IN
     Rowe, James Baber
PA
     Australia
S0
     PCT Int. Appl., 39 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
     ICM A61K031-445
     ICS A61K031-71; A61K033-06; A61K038-43; A61K038-46; A61K038-47
CC
     1-12 (Pharmacology)
     Section cross-reference(s): 17, 63
FAN. CNT 1
                                                                     DATE
     PATENT NO.
                          KIND
                                              APPLICATION NO.
                                 DATE
                                 19960711
                                              WO 1995-AU884
                                                                     19951229
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     WO 9620709
                           A1
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             NE,
                 SN,
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     CA 2208986
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                                 19960711
                                              CA 1995-2208986
                                                                     19951229
     AU 9643245
                                              AU 1996-43245
                                                                     19951229
                                 19960724
                           A1
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                                 19981105
                           B2
                                 19971015
                                              EP 1995-942004
                                                                     19951229
     EP 800394
                           A1
         R: AT, BE, CH,
                          DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT,
                          LV
                                                                     19951229
     NZ 297961
                           Α
                                 20000526
                                              NZ 1995-297961
                                              US 1997-860562
                                                                     19970829
     US 5985891
                                 19991116
                           Α
PRAI AU 1994-338
                                 19941229
     WO 1995-AU884
                                 19951229
CLASS
 PATENT NO.
                  CLASS
                         PATENT FAMILY CLASSIFICATION CODES
 WO 9620709
                  ICM
                         A61K031-445
                         A61K031-71; A61K033-06; A61K038-43; A61K038-46;
                  ICS
                         A61K038-47
                         A61K031/00+A; A61K031/424; A61K031/444; A61K038/46+M;
 WO 9620709
                 ECLA
                         A61K045/06
 US 5985891
                 ECLA
                         A61K031/00+A; A61K031/424; A61K031/444; A61K038/46+M;
                         A61K045/06
     This invention relates to a method for the treatment or prophylaxis of
AB
     adverse behavior, diarrhea, a skin disorder or an infection of the hind
     gut resulting from the accumulation of acid in the gastrointestinal tract
     of a human or an animal, said accumulation resulting from the fermentation of
     carbohydrate in the gastrointestinal tract of said human or animal, which
     method comprises administering to said human or animal an effective amount
     of an agent capable of preventing or controlling fermentative acidosis in
     the gastrointestinal tract.
     antidiarrheal digestive tract disorder behavior therapy
ST
IT
     Cereal
     Hav
        (digestion of; prevention of adverse behavior, diarrhea, skin disorders
        and infections of the hind gut associated with acidic conditions in humans
        and animals)
IT
     Galactosides
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (enzymes degrading; prevention of adverse behavior, diarrhea, skin
        disorders and infections of the hind gut associated with acidic conditions
        in humans and animals)
     Carbohydrates and Sugars, biological studies
     RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
        (fermentation of; prevention of adverse behavior, diarrhea, skin disorders
        and infections of the hind gut associated with acidic conditions in humans
        and animals)
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Qazi 10/642194

Page 47

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Diarrhea
        (inhibitors; prevention of adverse behavior, diarrhea, skin disorders
        and infections of the hind gut associated with acidic conditions in humans
        and animals)
IT
     Fermentation
        (of carbohydrates; prevention of adverse behavior, diarrhea, skin
        disorders and infections of the hind gut associated with acidic conditions
        in humans and animals)
ΙT
        (piglets; prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
ΙT
     Barley
     Canis familiaris
     Chicken
     Coprophagy
     Feeding experiment
     Horse
     Hyperkinesia
     Mental disorder
     Psoriasis
     Serpulina hyodysenteriae
     Sheep
     Skin, disease
        (prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
IΤ
     Polysaccharides, biological studies
     RL: ADV (Adverse effect, including toxicity); BPR (Biological process);
     BSU (Biological study, unclassified); BIOL (Biological study); PROC
     (Process)
        (prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
     Clays, biological studies
     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
     process); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
     Antibiotics
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
IΤ
     Bentonite, biological studies
     Zeolite-group minerals
     RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
        (prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
ΙT
     Sulfonamides
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
IT
        (swine; prevention of adverse behavior, diarrhea, skin disorders and
        infections of the hind gut associated with acidic conditions in humans and
        animals)
IT
     Antibiotics
        (aminoglycoside, prevention of adverse behavior, diarrhea, skin
        disorders and infections of the hind gut associated with acidic conditions
        in humans and animals)
     Mental disorder
IT
        (attention deficit, prevention of adverse behavior, diarrhea, skin
```

Qazi 10/642194

Page 48

disorders and infections of the hind gut associated with acidic conditions in humans and animals)

IT Intestine

(cecum, prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

IΤ Digestive tract

(disease, prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

IT Digestive tract

(disease, indigestion, acid accumulation in; prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

ΙT Behavior

(disorder, prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

IT Bacteria

> (gram-pos., prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

Digestive tract ΙT

(hindgut, disorders; prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

ΙT Bacteria

(intestinal, lactic acid production by; prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

ΙT Intestine, disease

(large, irritable bowel disease; prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

IT Pharmaceutical dosage forms

(oral, prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

IT

50-21-5, Lactic acid, biological studies RL: ADV (Adverse effect, including toxicity); BSU (Biological study, unclassified); MFM (Metabolic formation); BIOL (Biological study); FORM (Formation, nonpreparative)

(bacterial production of; prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

512-69-6 9000-69-5, Pectins 9037-55-2, Galacturonan 9060-75-7, L-Arabinan 11078-30-1, 9037-55-2, Galactan

39280-21-2, Galactomannan 37294-28-3, Xyloglucan

Rhamnogalacturonan

RL: BSU (Biological study, unclassified); BIOL (Biological study) (enzymes degrading; prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

IT 9040-13-5, Virginiamycin m

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES

(prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

1318-74-7, Kaolinite, biological studies 1318-93-0, Montmorillonite RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses) (prevention of adverse behavior, diarrhea, skin disorders and infections of the hind gut associated with acidic conditions in humans and animals)

60-54-8, Tetracycline 114-07-8, Erythromycin 804-36-4, Nitrovin 1393-48-2, Thiostrepton 1393-68-6, Bottromycin 1405-89-6, Bacitracin zinc 1406-05-9D, Penicillin, derivs. 1476-53-5, Novobiocin sodium 1695-77-8, Spectinomycin 9000-92-4, Amylase 9001-22-3, Emulsin

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9001-42-7, Maltase 9001-57-4, Invertase
                                                      9015-78-5, Glucanase
     9025-35-8, \alpha-Galactosidase 9032-08-0, Amyloglucosidase
     9074-98-0, β-Glucanase 11006-76-1, Streptogramin 11015-37-5,
     Flavomycin
                   11017-43-9, Siomycin 11054-70-9, Lasalocid
     11111-12-9D, Cephalosporin, derivs.
     11115-82-5, Enramycin 12609-84-6, Thiopeptin 13721-01-2D, derivs. 37244-77-2, Sporangiomycin 37278-89-0, Xylanase 37332-99-3, Avoparcin
     53003-10-4, Salinomycin 55134-13-9, Narasin 55297-95-5, Tiamulin
     55852-84-1, Bacitracin methylene disalicylate 65454-59-3, Sulfomycin 75139-06-9, Tetronasi 117742-13-9, Ardacin
                                                           65454-16-2, Taitomycin
                                 75139-06-9, Tetronasin
                                                             80738-43-8, Lincosamide
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
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         infections of the hind gut associated with acidic conditions in humans and
         animals)
IT
     11078-30-1, Galactomannan
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (enzymes degrading; prevention of adverse behavior, diarrhea, skin
         disorders and infections of the hind gut associated with acidic conditions
         in humans and animals)
     11078-30-1 HCAPLUS
RN
     D-Galacto-D-mannan (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
IT
     11111-12-9D, Cephalosporin, derivs.
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (prevention of adverse behavior, diarrhea, skin disorders and
         infections of the hind gut associated with acidic conditions in humans and
         animals)
RN
     11111-12-9 HCAPLUS
     Cephalosporin (9CI)
                             (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
L44 ANSWER 11 OF 12 HCAPLUS COPYRIGHT 2005 ACS on STN
     1995:842635 HCAPLUS
AN
DN
     123:237834
     Entered STN: 10 Oct 1995
ED
     Transdermal patch for release of active agents from hot-melt adhesives
ΤI
     Hoffmann, Hans-Rainer; Roreger, Michael
IN
     ITs Lohmann Therapie-Systeme GmbH und Co. KG, Germany
PA
     Ger., 14 pp.
     CODEN: GWXXAW
DT
     Patent
LA
     German
     ICM A61L015-44
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FAN. CNT 1
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                                                                           19940513
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A1 19951205 AU 1995-26127 19950508
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     HU 75288
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NO 9604799
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     GR 3035478
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                                   20010531
                                                GR 2001-400320
                                                                         20010228
PRAI DE 1994-4416927
                                   19940513
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     WO 1995-EP1724
                                   19950508
CLASS
 PATENT NO.
                          PATENT FAMILY CLASSIFICATION CODES
                  CLASS
 DE 4416927
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                          A61L015-44
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 US 6190689
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                          A01N025/24; A61K009/70E; A61L015/44; A61L015/58;
                          C09J011/00
     A device for release of active agents from a hot-melt adhesive, with
AB
     uniform or nonuniform distribution of the agent in the adhesive, comprises
     a hydrophilic adhesive containing ≥1 water-soluble or water-swellable
     polymer [e.g. gelatin, Na alginate, pectin, starch (derivative), PVP, PEG],
     ≥1 water-soluble, fusible adhesive resin [e.g. pantothenol, honey,
     low-mol.-weight sugar (derivative)], and the active agent. The processing temperature
     for the adhesive is 40-150°, preferably 80-120°. Thus, a melt containing hydroabietyl alc. 25, ethylene/vinyl acetate (72:28) copolymer 10, C4-5 polyalkadiene 10, and acetylsalicylic acid 10 parts was prepared at
     120^{\circ} , cooled to 90^{\circ} , mixed with a melt containing pantothenol
     25, Na alginate 10, and gelatin 10 parts, and spread on siliconized paper
     composite to a thickness of 300 g/m2.
     transdermal patch hot melt adhesive
ĪΤ
     Resins
     RL: DEV (Device component use); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
         (adhesive; transdermal patch for release of active agents from hot-melt
        adhesives)
IT
     Chamomile
     Valerian
         (extract; transdermal patch for release of active agents from hot-melt
        adhesives)
IT
     Plant.
         (exts.; transdermal patch for release of active agents from hot-melt
        adhesives)
     Steroids, biological studies RL: BAC (Biological activity or effector, except adverse); BSU (Biological
IT
     study, unclassified); DEV (Device component use); THU (Therapeutic use);
     BIOL (Biological study); USES (Uses)
         (hormones; transdermal patch for release of active agents from hot-melt
        adhesives)
IT
     Tobacco
         (powdered; transdermal patch for release of active agents from hot-melt
        adhesives)
IT
     Arnica
     Marigold
     St. -John's-wort
         (tincture; transdermal patch for release of active agents from hot-melt
        adhesives)
IT
     Cosmetics
     Echinacea angustifolia
     Fucus vesiculosus
     Honey
     Hop
     Veterinary medicine
     Wound healing promoters
         (transdermal patch for release of active agents from hot-melt
        adhesives)
     Chlorophylls, biological studies
     RL: BAC (Biological activity or effector, except adverse); BSU (Biological
     study, unclassified); DEV (Device component use); THU (Therapeutic use);
     BIOL (Biological study); USES (Uses)
```

(transdermal patch for release of active agents from hot-melt adhesives)

ΙT Carbohydrates and Sugars, biological studies

Collagens, biological studies

Gelatins, biological studies

RL: DEV (Device component use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(transdermal patch for release of active agents from hot-melt adhesives)

Polymers, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (water-soluble/-swellable; transdermal patch for release of active agents from hot-melt adhesives)

Essential oils

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); DEV (Device component use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(chamomile, German, transdermal patch for release of active agents from hot-melt adhesives)

Essential oils

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); DEV (Device component use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(eucalyptus, transdermal patch for release of active agents from

hot-melt adhesives)

Tea products

(green, transdermal patch for release of active agents from hot-melt adhesives)

ΙT Adhesives

> (hot-melt, transdermal patch for release of active agents from hot-melt adhesives)

ΙT Essential oils

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); DEV (Device component use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(pine leaf, transdermal patch for release of active agents from

hot-melt adhesives)

Alkadienes IT

RL: DEV (Device component use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(polymers, transdermal patch for release of active agents from hot-melt adhesives)

IT Pharmaceutical dosage forms

(transdermal, transdermal patch for release of active agents from

hot-melt adhesives) 50-78-2, Acetylsalicylic acid 50-28-2, Estradiol, biological studies 69-72-7, Salicylic acid, biological studies 137-58-6D, Lidocaine, eutectic with prilocaine 721-50-6D, Prilocaine, eutectic with lidocaine 1406-18-4, Vitamin E 18472-51-0, Chlorhexidine digluconate RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); DEV (Device component use); THU (Therapeutic use);

BIOL (Biological study); USES (Uses) (transdermal patch for release of active agents from hot-melt

adhesives)

50-99-7, Glucose, biological studies 56-81-5, Gl studies 57-48-7, D-Fructose, biological studies 56-81-5, Glycerin, biological IT 57-50-1, Sucrose, biological studies 81-13-0, Pantothenol 126-13-6, Sucrose acetate 9000-07-1, Carrageenan 9000-69-5, Pectin 9002-89-5, 9004-32-4, Sodium CM-cellulose 9002-89-5, isobutyrate 9003-39-8, PVP Poly(vinyl alcohol) 9004-57-3, Ethylcellulose 9004-34-6D, Cellulose, derivs. 9005-25-8, Starch, biological studies 9005-25-8D, Starch, derivs. 9005-38-3, 9063-38-1, Sodium carboxymethylstarch 11078-30-1 Sodium alginate , Galactomannan 11078-30-1D, Galactomannan,

11138-66-2, Xanthan 24937-78-8, Ethylene/vinyl acetate derivs. 25086-89-9, Vinylpyrrolidone/vinyl acetate copolymer 25322-68-3, PEG 25322-69-4, Polypropylene glycol 26266-77-3, Dihydroabietyl alcohol

RL: DEV (Device component use); THU (Therapeutic use); BIOL (Biological

study); USES (Uses)

(transdermal patch for release of active agents from hot-melt

```
adhesives)
IT
     11078-30-1, Galactomannan 11078-30-1D,
     Galactomannan, derivs.
     RL: DEV (Device component use); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
         (transdermal patch for release of active agents from hot-melt
        adhesives)
     11078-30-1 HCAPLUS
RN
CN
     D-Galacto-D-mannan (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
RN
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     D-Galacto-D-mannan (9CI)
CN
                                 (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
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     1977:411647 HCAPLUS
AN
DN
     87:11647
ED
     Entered STN: 12 May 1984
ΤI
     Controlled release tablets
     Walton, Richard Wiegmann; De Lorimier, Albert Emmanuel
IN
PA
     E. R. Squibb and Sons, Inc., USA
     Ger. Offen., 19 pp.
     CODEN: GWXXBX
DT
     Patent
LA
     German
     A61K009-26
     63-6 (Pharmaceuticals)
CC
FAN. CNT 1
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PRAI US 1975-621316
                            Α
                                  19751010
CLASS
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                  IC
                         A61K009-26
 DE 2645547
     Controlled release tablets comprise a matrix of a drug dispersed in a
     mixture of polyvinylpyrrolidone (PVP) [9003-39-8] and a
     hydrophilic carboxy group-containing vinyl polymer coated with an H2O-insol.,
     H2O-permeable mixture of a moderately H2O-soluble hydrophobic polymer and an
     H2O-soluble hydrophilic polymer. The coating breaks open when the matrix
     swells. For example, 790 mg tablets were compressed from a mixture of procaineamide-HCl [614-39-1] 500, PVP 144, carbopol 934 [9007-16-3] 96,
     microcryst. cellulose 23.4, powdered carnauba wax 15.6, stearic acid 7.8, and
     silicic acid 3.95 g. The tablets were coated with a 0.09-0.1 mm layer of
     a mixture of hydroxypropyl methyl cellulose [9004-65-3] 30, ethyl cellulose
     [9004-57-3] 20 and triethyl citrate 2 g/L, 284.4 mL 99% iso-PrOH and 664
     mL CH2C12. In H2O at 37^\circ , the tablets released 14.6, 20.1, 13.3, 20.0, 1.5, and 2.1% of the initial drug content during h 1, 2, 3, 4, 5 and
     8, resp., compared to 40.0, 11.8, 14.4, 9.8, 3.6 and 0% from uncoated
```

Tablet controlled release; procainamide controlled release tablet; cephadrin controlled release tablet

tablets.

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IT
      Tablets
          (controlled-release, vinyl polymers in)
      614-39-1 38821-53-3 RL: BIOL (Biological study)
          (controlled-release tablet)
IT
      9004-57-3 9004-65-3 9050-31-1
      RL: BIOL (Biological study)
          (in controlled release tablet coating)
      9003-39-8 9007-16-3
ΙT
      RL: BIOL (Biological study)
      (in controlled-release tablet matrix) 38821-53-3
IT
      RL: BIOL (Biological study)
          (controlled-release tablet)
      38821-53-3 HCAPLUS
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-amino-1,4-cyclohexadien-1-ylacetyl]amino]-3-methyl-8-oxo-,
(6R,7R)- (9CI) (CA INDEX NAME)
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Absolute stereochemistry.

=> b home FILE 'HOME' ENTERED AT 16:31:22 ON 15 APR 2005 => b reg FILE 'REGISTRY' ENTERED AT 08:03:24 ON 19 APR 2005 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2005 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 18 APR 2005 HIGHEST RN 848724-42-5 DICTIONARY FILE UPDATES: 18 APR 2005 HIGHEST RN 848724-42-5

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 18, 2005

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

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L29 ANSWER 1 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 770660-21-4 REGISTRY
ED Entered STN: 27 Oct 2004
CN 5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid,
7-[[(2-amino-4-thiazolyl) (methoxyimino)acetyl]amino]-8-oxo-,
[6R-(6α, 7β)]- (9CI) (CA INDEX NAME)
FS STEREOSEARCH
MF C13 H13 N5 05 S2
CI COM
SR CA

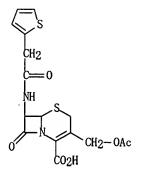
Absolute stereochemistry. Double bond geometry unknown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

L29 ANSWER 2 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

Absolute stereochemistry.

#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*



# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

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L29 ANSWER 4 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN 442852-34-8 REGISTRY
ED Entered STN: 07 Aug 2002
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
    7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino)acetyl]amino]-3-[(1Z)-2-(4-methyl-5-thiazolyl)ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy)methyl ester, dihydrochloride, (6R, 7R)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH
MF C25 H28 N6 07 S3 . 2 C1 H

SR CA
LC STN Files: CA, CAPLUS
CRN (117467-28-4)
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Absolute stereochemistry. Double bond geometry as shown.

#### ●2 HC1

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 5 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

ED

ANSWER 5 OF 118 REGISTRY COPYRIGHT 2005 ACS ON SIN

442852-33-7 REGISTRY
Entered STN: 07 Aug 2002

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-3-[(1Z)-2-(4-methyl-5-thiazolyl) ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy) methyl ester, (6R, 7R)-, sulfate (1:1) (9CI) (CA INDEX NAME)

STEREOSEARCH

MF C25 H28 N6 O7 S3 . H2 O4 S

SR CA

STN Files: CA, CAPLUS

> CM 1

117467-28-4 CRN C25 H28 N6 O7 S3 CMF

Absolute stereochemistry. Double bond geometry as shown.

CM 2

CRN 7664-93-9 CMF H2 04 S

### 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 6 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN 442852-32-6 REGISTRY
ED Entered STN: 07 Aug 2002

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-3-[(1Z)-2-(4-methyl-5-thiazolyl) ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy) methyl
ester, (6R, 7R)-, (2R, 3R)-2, 3-dihydroxybutanedioate (1:1) (9CI) (CA INDEX NAME)

**STEREOSEARCH** 

MF C25 H28 N6 O7 S3 . C4 H6 O6

SR

STN Files: CA, CAPLUS

CM

CRN 117467-28-4 CMF C25 H28 N6 O7 S3

Absolute stereochemistry. Double bond geometry as shown.

CM

CRN 87-69-4 CMF C4 H6 O6

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

- L29
- RN
- ED

ANSWER 7 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 442852-31-5 REGISTRY Entered STN: 07 Aug 2002 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7 [5](07) 7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl] amino]-3-[(1Z)-2-(4-methoxyimino)] methyl-5-thiazolyl)ethenyl]-8-oxo-, (2,2-dimethyl-1-oxopropoxy)methyl ester, (6R,7R)-, ethanedioate (1:1) (9CI) (CA INDEX NAME)

FS STEREOSEARCH
MF C25 H28 N6 07 S3 . C2 H2 O4
SR CA
LC STN Files: CA, CAPLUS

CM 1

CRN 117467-28-4
CMF C25 H28 N6 07 S3

Absolute stereochemistry.

Double bond geometry as shown.

CM 2

CRN 144-62-7 CMF C2 H2 O4

HO-C-C-OH

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 8 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 442852-30-4 REGISTRY
ED Entered STN: 07 Aug 2002
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino)acetyl]amino]-3-[(1Z)-2-(4-methyl-5-thiazolyl)ethenyl]-8-oxo-, (2,2-dimethyl-1-oxopropoxy)methyl ester, (6R, 7R)-, diethanesulfonate (9CI) (CA INDEX NAME)
FS STEREOSEARCH
MF C25 H28 N6 07 S3 . 2 C2 H6 O3 S
SR CA
LC STN Files: CA, CAPLUS

CM 1

Absolute stereochemistry.

Double bond geometry as shown.

117467-28-4

C25 H28 N6 O7 S3

CRN

CMF

CM

CRN 594-45-6 CMF C2 H6 O3 S

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29

RN

ED

ANSWER 9 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 442852-29-1 REGISTRY Entered STN: 07 Aug 2002 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-3-[(1Z)-2-(4-methyl-5-thiazolyl) ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy) methyl ester, (6R, 7R)-, dimethanesulfonate (9CI) (CA INDEX NAME) CN

FS STEREOSEARCH

C25 H28 N6 O7 S3 . 2 C H4 O3 S MF

SR LC CA

STN Files: CA, CAPLUS

CM

CRN 117467-28-4 C25 H28 N6 O7 S3 CMF

Absolute stereochemistry. Double bond geometry as shown.

CM

CRN 75-75-2 C H4 03 S

## 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 10 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 352467-09-5 REGISTRY Entered STN: 23 Aug 2001

RN

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, (6R, 7R)-, mononitrate (salt) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Cefadroxil nitrate

STEREOSEARCH FS

C16 H17 N3 O5 S . H N O3

SR

STN Files: CA, CAPLUS, USPAT2, USPATFULL

CM

50370-12-2 CRN C16 H17 N3 05 S **CMF** 

Absolute stereochemistry.

$$\begin{array}{c|c} & & & \\ &$$

CM 2

CRN 7697-37-2 H N 03 CMF

### 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 11 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

ED

352465-77-1 REGISTRY
Entered STN: 23 Aug 2001
5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-, CN (6R, 7R)-, nitrate (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Ceftizoxime nitrate

FS STEREOSEARCH

MF C13 H13 N5 O5 S2 . x H N O3

SR CA

STN Files: CA, CAPLUS, USPAT2, USPATFULL

CM

CRN 68401-81-0 CMF C13 H13 N5 O5 S2

Absolute stereochemistry. Double bond geometry as shown.

CM

CRN 7697-37-2 CMF H N 03

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 12 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

ED

352465-69-1 REGISTRY
Entered STN: 23 Aug 2001
5-Thia-1-azabicyclo[4, 2, 0]oct-2-ene-2-carboxylic acid, 7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino)acetyl]amino]-3(methoxymethyl)-8-oxo-, 1-[[(1-methylethoxy)carbonyl]oxy]ethyl ester,
(6R, 7R)-, nitrate (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Cefpodoxime proxetil nitrate

FS STEREOSEARCH

MF C21 H27 N5 O9 S2 . x H N O3

SR

STN Files: CA, CAPLUS, USPAT2, USPATFULL

CM 1

CRN 87239-81-4 CMF C21 H27 N5 09 S2

Absolute stereochemistry. Double bond geometry as shown.

CM

CRN 7697-37-2 CMF H N 03

# 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 13 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 352465-61-3 REGISTRY
ED Entered STN: 23 Aug 2001
CN 15-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-hydroxyphenylacety1]amino]-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, (6R, 7R)-, mononitrate (salt) (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Cefamandole nitrate

FS STEREOSEARCH

MF C18 H18 N6 O5 S2 . H N O3

SR CA

STN Files: CA, CAPLUS, USPAT2, USPATFULL

CM

34444-01-4 CMF C18 H18 N6 O5 S2

Absolute stereochemistry.

CM

CRN 7697-37-2

CMF H N 03

# 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

RN

ED

ANSWER 14 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 352465-55-5 REGISTRY
Entered STN: 23 Aug 2001
5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1-propenyl)-,
(6R, 7R)-, mononitrate (salt) (9CI) (CA INDEX NAME)
FR NAMES: CN

OTHER NAMES:

CN Cefprozil nitrate

FS STEREOSEARCH

MF C18 H19 N3 O5 S . H N O3

SR

STN Files: CA, CAPLUS, USPAT2, USPATFULL

CM

92665-29-7 CMF C18 H19 N3 O5 S

Absolute stereochemistry. Double bond geometry unknown.

CM

CRN 7697-37-2 CMF H N 03

# 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 15 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 352465-52-2 REGISTRY ED Entered STN: 23 Aug 2001

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, (6R,7R)-, mononitrate CN (9CI) (CA INDEX NAME)
OTHER NAMES:

Cefaclor nitrate CN

FS STEREOSEARCH

MF C15 H14 C1 N3 O4 S . H N O3

SR CA

STN Files: CA, CAPLUS, USPAT7, USPATFULL CM

CRN 53994-73-3 C15 H14 C1 N3 O4 S

Absolute stereochemistry.

CM

7697-37-2 CRN H N 03 CMF

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 16 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

ED

ANSWER TO OF THE REGISTRY COPYRIGHT 2005 ACS ON SIN

350230-49-8 REGISTRY
Entered STN: 03 Aug 2001
5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, dihydrate,
(6R, 7R)- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Cephalothin dihydrate

FS STEREOSEARCH

MF C16 H16 N2 O6 S2 . 2 H2 O

SR CA

CA, CAPLUS, TOXCENTER, USPAT2, USPATFULL LC STN Files:

CRN (153-61-7)

Absolute stereochemistry.

●2 H<sub>2</sub>0

4 REFERENCES IN FILE CA (1907 TO DATE) 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 17 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 190181-58-9 REGISTRY ED Entered STN: 20 Jun 1997

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[(hydroxyphenylacetyl)amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-5-yl]thio]methyl]-, disodium salt, [6R-(6 $\alpha$ , 7 $\beta$ )]-[partial]- (9CI) (CA INDEX NAME)

STEREOSEARCH

MF C18 H18 N6 O8 S3 . 2 Na

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

CRN (69088-92-2)

Absolute stereochemistry.

## Na

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 18 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 155723-35-6 REGISTRY ED Entered STN: 15 Jun 1994

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-3-[2-(3-methyl-4-isothiazolyl) ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy) methyl ester, [6R-[3(Z), 6 $\alpha$ , 7 $\beta$ (Z)]]- (9CI) (CA INDEX NAME)

STEREOSEARCH

MF C25 H28 N6 O7 S3

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry. Double bond geometry as shown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 19 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

ED

148774-47-4 REGISTRY
Entered STN: 20 Jul 1993
5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, CN 7-[[(2-amino-4-thiazoly1) (methoxyimino) acetyl]amino]-3-[2-(4-methyl-5-thiazoly1) ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy) methyl ester, [6R-[3(Z), 6 $\alpha$ , 7 $\beta$ (Z)]]- (9CI) (CA INDEX NAME)

STEREOSEARCH FS

C25 H28 N6 O7 S3 MF

SR CA

LC STN Files: CA, CAPLUS

Absolute stereochemistry. Double bond geometry as shown.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 20 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN 144790-28-3 REGISTRY

ED Entered STN: 08 Dec 1992

7-[[(2S)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, (6R, 7R)-(9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-,  $[6R-[6\alpha, 7\beta(S*)]]-$ 

OTHER NAMES:

L-Cefadroxil

FS STEREOSEARCH

MF C16 H17 N3 O5 S

SR CA

BEILSTEIN\*, CA, CAPLUS STN Files:

(\*File contains numerically searchable property data)

Absolute stereochemistry.

### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1907 TO DATE) 5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 21 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 138514-32-6 REGISTRY ED Entered STN: 24 Jan 1992

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl] amino]-3-[2-(4-methyl-5-thiazolyl) ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy) methyl ester,  $[6R-[3(E), 6\alpha, 7\beta(Z)]]-(9CI)$  (CA INDEX NAME)

OTHER NAMES:

ME 1207E CN

FS STEREOSEARCH

MF C25 H28 N6 07 S3

SR CA

LC BEILSTEIN\*, CA, CAPLUS STN Files:

(\*File contains numerically searchable property data)

Absolute stereochemistry. Double bond geometry as shown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 22 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 132850-29-4 REGISTRY Entered STN: 29 Mar 1991 L29

RN

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-, monosodium salt, hydrate, [6R-[6 $\alpha$ ,7 $\beta$ (Z)]]- (9CI) (CA INDEX NAME) STEREOSEARCH

MF C13 H13 N5 O5 S2 . x H2 O . Na

SR CA

LC STN Files: CA, CAPLUS, IMSPATENTS

(68401-81-0)

# 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 23 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 123458-61-7 REGISTRY ED Entered STN: 27 Oct 1989  $\_$ 5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-[[(2Z)-2-furanyl(methoxyimino)acetyl]amin CN o]-8-oxo-, 1-(acetyloxy)ethyl ester, (6R, 7R)- (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 3-[[(aminocarbonyl)oxy]methyl]-7-[[2-furanyl(methoxyimino)acetyl]amino]-8oxo-, 1-(acetyloxy)ethyl ester,  $[6R-[6\alpha, 7\beta(Z)]]$ -OTHER NAMES: △2-Cefuroxime axetil CN **STEREOSEARCH** FS MF C20 H22 N4 010 S SR CA STN Files: CA, CAPLUS, CASREACT, USPATFULL

Absolute stereochemistry.

Double bond geometry as shown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1907 TO DATE) 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 24 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 121412-77-9 REGISTRY Entered STN: 30 Jun 1989 RN ED CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1Z)-1-propenyl-, (6R, 7R)- (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1-propenyl)-,  $[6R-[3(Z), 6\alpha, 7\beta(R*)]]-$ OTHER NAMES: cis-Cefprozil CN FS STEREOSEARCH DR 136427-55-9 C18 H19 N3 O5 S MF CI COM SR CA LC CA, CAPLUS, CASREACT, IMSPATENTS, IMSRESEARCH, MRCK\*, STN Files: USPATFULL (\*File contains numerically searchable property data)

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

10 REFERENCES IN FILE CA (1907 TO DATE) 10 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 25 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 121123-17-9 REGISTRY Entered STN: 16 Jun 1989 L29

RN

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,

7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1-propenyl)-, monohydrate, (6R, 7R)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,

7-[[amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1-propenyl)-, monohydrate, [6R-[6 $\alpha$ , 7 $\beta$ (R\*)]]-

OTHER NAMES:

Cefprozil monohydrate CN

FS STEREOSEARCH

MF C18 H19 N3 O5 S . H2 O

CAS Client Services

N Files: CA, CAPLUS, IMSPATENTS, IMSRESEARCH, MRCK\*, PROUSDDR, PS, SYNTHLINE, TOXCENTER, USAN, USPATFULL STN Files:

(\*File contains numerically searchable property data)

CRN (92665-29-7)

Absolute stereochemistry. Double bond geometry unknown.

$$\begin{array}{c|c} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\$$

● H20

3 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 26 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

119922-85-9 REGISTRY Entered STN: 07 Apr 1989 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, hydrate (2:1), (6R, 7R) - (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, hydrate (2:1),

Page 17 Qazi 10/642194

```
[6R-[6\alpha, 7\beta(R*)]]-
OTHER NAMES:
     Cefadroxil hemihydrate
FS
     STEREOSEARCH
     C16 H17 N3 O5 S . 1/2 H2 O
MF
SR
     STN Files:
                  BEILSTEIN*, CA, CAPLUS, CIN, DIOGENES, IMSPATENTS, USAN,
       USPATFULL
         (*File contains numerically searchable property data)
     (50370-12-2)
```

Absolute stereochemistry.

$$\begin{array}{c|c} & & & \\ &$$

## ●1/2 H<sub>2</sub>0

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3 REFERENCES IN FILE CA (1907 TO DATE)
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3 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L29 ANSWER 27 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
        117467-28-4 REGISTRY
RN
        Entered STN: 11 Nov 1988
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-3-[(1Z)-2-(4-methyl-5-thiazolyl) ethenyl]-8-oxo-, (2, 2-dimethyl-1-oxopropoxy) methyl ester, (6R, 7R)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-3-[2-(4-methyl-5-
        thiazolyl)ethenyl]-8-oxo-, (2,2-dimethyl-1-oxopropoxy)methyl ester,
        [6R-[3(Z), 6\alpha, 7\beta(Z)]]-
OTHER NAMES:
        CCRIS 7768
CN
       Cefditoren pivaloyloxymethyl ester
CN
        Cefditoren pivoxil
       Cefditoren pivoxyl
CN
       Cefditorin
CN
CN
CN
       ME 1207
        Meiact
        Spectracef
FS
        STEREOSEARCH
        C25 H28 N6 07 S3
MF
CI
       COM
SR
           IN Files: ADISINSIGHT, ADISNEWS, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CIN, DDFU, DIOGENES, DRUGU, EMBASE, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*, PHAR, PROMT, PROUSDDR, PS, RTECS*, SYNTHLINE, TOXCENTER, USAN,
        STN Files:
           USPAT2, USPATFULL
               (*File contains numerically searchable property data)
```

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

123 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
124 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 28 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

117091-43-7 REGISTRY Entered STN: 22 Oct 1988 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyl-113C-oxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-trans)-(9CI) (CA INDEX NAME)

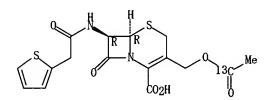
**STEREOSEARCH** 

MF C16 H16 N2 O6 S2

SR CA

STN Files: CA, CAPLUS

Absolute stereochemistry.



1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

- L29 ANSWER 29 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
- RN
- 109323-67-3 REGISTRY Entered STN: 18 Jul 1987 ED
- 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-, monohydrochloride, monohydrate, [6R-[6 $\alpha$ , 7 $\beta$ (2)]]- (9CI) (CA INDEX NAME)
- STEREOSEARCH
- C13 H13 N5 05 S2 . C1 H . H2 0
- SR
- BEILSTEIN\*, CA, CAPLUS, IMSPATENTS

(\*File contains numerically searchable property data)

(68401 - 81 - 0)CRN

HC1

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 30 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN

104145-87-1 REGISTRY Entered STN: 13 Sep 1986 ED

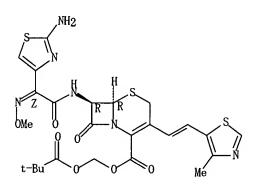
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino)acetyl]amino]-3-[2-(4-methyl-5thiazolyl)ethenyl]-8-oxo-, (2,2-dimethyl-1-oxopropoxy)methyl ester, [6R-[6\alpha, 7\beta(Z)]]- (9CI) (CA INDEX NAME)
STEREOSEARCH

MF C25 H28 N6 07 S3

SR CA

STN Files: LC BEILSTEIN\*, CA, CAPLUS, IMSPATENTS, IMSRESEARCH, PS, USPATFULL (\*File contains numerically searchable property data)

Absolute stereochemistry. Double bond geometry as described by E or Z.



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 31 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

ED

102044-69-9 REGISTRY
Entered STN: 10 May 1986
5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, CN 7-[[(2-amino-4-thiazolyl)(methoxyimino)acetyl]amino]-8-oxo-,  $[2R-[2\alpha, 6\alpha, 7\beta(Z)]]-(9CI)$  (CA INDEX NAME)

FS STEREOSEARCH

C13 H13 N5 O5 S2 MF

SR LC STN Files: BEILSTEIN\*, CA, CAPLUS, CASREACT (\*File contains numerically searchable property data)

Absolute stereochemistry. Double bond geometry as shown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 32 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 97164-53-9 REGISTRY Entered STN: 13 Jul 1985 L29

RN

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino)acetyl]amino]-8-oxo-, monosodium salt, [6R-[6 $\alpha$ , 7 $\beta$ (E)]]- (9CI) (CA INDEX NAME)

STEREOSEARCH

MF

C13 H13 N5 O5 S2 . Na STN Files: BEILSTEIN\*, CA, CAPLUS, USPATFULL LC (\*File contains numerically searchable property data)

CRN (68403-31-6)

Absolute stereochemistry. Double bond geometry as shown.

Na

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 33 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**94539-08-9** REGISTRY Entered STN: 03 Feb 1985 ED

5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(3-thienyl)acetamido]-, acetate (7CI)

INDEX NAME)
FS STEREOSEARCH
MF C16 H16 N2 O6 S2

LC STN Files: CAOLD

Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

#### 1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

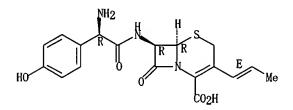
L29 ANSWER 34 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 92676-86-3 REGISTRY RN Entered STN: 17 Dec 1984 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, ED CN 7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1E)-1-propenyl-, (6R, 7R) - (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1-propenyl)-,  $[6R-[3(E), 6\alpha, 7\beta(R*)]]-$ OTHER NAMES: BMY 28167 CN CN trans-Cefprozil FS STEREOSEARCH DR 136427-56-0 C18 H19 N3 O5 S MF CI COM CA, CAPLUS, IMSPATENTS, IMSRESEARCH, MRCK\*, TOXCENTER, LC STN Files:

(\*File contains numerically searchable property data)

Absolute stereochemistry.

Double bond geometry as shown.

USPATFULL



# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

13 REFERENCES IN FILE CA (1907 TO DATE)
13 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 35 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 92665-29-7 REGISTRY ED Entered STN: 17 Dec 1984

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1-propenyl)-,
(6R, 7R)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

Page 22

```
5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
        7-[[amino(4-hydroxyphenyl)acetyl]amino]-8-oxo-3-(1-propenyl)-,
        [6R-[6\alpha, 7\beta(R*)]]-
OTHER NAMES:
        BMY 28100
CN
        Cefprozil
CN
        Cefprozile
CN
        Cefzil
CN
        Procef
CN
FS
        Prozef
        STEREOSEARCH
MF
        C18 H19 N3 O5 S
CI
        COM
           OM
TN Files: ADISINSIGHT, ADISNEWS, ANABSTR, BIOBUSINESS, BIOSIS,
BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CIN,
DDFU, DIOGENES, DRUGU, EMBASE, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH,
IPA, MEDLINE, MRCK*, PHAR, PROMT, PROUSDDR, PS, RTECS*, SYNTHLINE,
TOXCENTER, USAN, USPAT2, USPATFULL
        STN Files:
               (*File contains numerically searchable property data)
        Other Sources:
                                    WHO
```

Absolute stereochemistry. Double bond geometry unknown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

```
7 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
             311 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L29 ANSWER 36 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
     91828-55-6 REGISTRY
     Entered STN: 16 Nov 1984
ED
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
     7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, dihydrate, (6R, 7R)-
     (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
     7-[(aminophenylacetyl)amino]-3-chloro-8-oxo-, dihydrate,
     [6R-[6\alpha, 7\beta(R*)]]-
OTHER NAMES:
     Cefaclor dihydrate
CN
FS
     STEREOSEARCH
     C15 H14 C1 N3 O4 S . 2 H2 O
STN Files: BIOSIS, CA, CAPLUS, IMSPATENTS, IPA, USPATFULL
MF
LC
     (53994 - 73 - 3)
```

309 REFERENCES IN FILE CA (1907 TO DATE)

Qazi 10/642194

# ●2 H<sub>2</sub>0

```
3 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
3 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L29 ANSWER 37 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 87239-81-4 REGISTRY ED Entered STN: 16 Nov 1984
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-3-
(methoxymethyl)-8-oxo-, 1-[[(1-methylethoxy) carbonyl]oxy]ethyl ester,
(6R, 7R)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl)(methoxyimino)acetyl]amino]-3-(methoxymethyl)-8-
       oxo-, 1-[[(1-methylethoxy)carbonyl]oxy]ethyl ester, [6R-[6α, 7β(Z)]]-
OTHER NAMES:
        (RS)-1-(Isopropoxycarbonyloxy) ethyl (+)-(6R,7R)-7-[2-(2-amino-4-thiazolyl)-1-(1sopropoxycarbonyloxy)]
        2-[(Z)-(methoxyimino]acetamido]-3-methoxymethyl-8-oxo-5-thia-1-
       azabicyclo[4.2.0]oct-2-ene-2-carboxylate
       Antibiotic CS 807
CN
       Banan
CN
       Cefodox
CN
       Cefpodoxime proxetil
CN
       Cephpodoxime proxetil
       CS 807
       CS 807 (pharmaceutical)
CN
       0relox
CN
       Otreon
       RU 51807
CN
       U 76252
CN
       Vantin
FS
       STEREOSEARCH
DR
       95242-58-3
MF
       C21 H27 N5 O9 S2
CI
       COM
                           ADISNEWS, AGRICOLA, ANABSTR, BIOBUSINESS, BIOSIS, BIOTECHNO,
          CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CHEMCATS, CIN, DIOGENES, EMBASE, IMSCOSEARCH, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*, PATDPASPC, PHAR, PROMT, PROUSDDR, RTECS*, SYNTHLINE, TOXCENTER, USAN,
           USPAT2, USPATFULL
              (*File contains numerically searchable property data)
```

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

253 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 254 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 38 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 87018-84-6 REGISTRY Entered STN: 16 Nov 1984

RN

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2-amino-4-thiazolyl) (methoxylmino) acetyl]amino]-8-oxo-, monopotassium CNsalt,  $[6R-[6\alpha,7\beta(Z)]]-(9CI)$  (CA INDEX NAME) STEREOSEARCH

MF C13 H13 N5 O5 S2 . K

CA, CAPLUS, IMSPATENTS LC STN Files:

CRN (68401-81-0)

Absolute stereochemistry. Double bond geometry as shown.

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

- ANSWER 39 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
- RN
- 86068-04-4 REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(thienylacetyl)amino]-, monosodium salt, (6R-trans)- (9CI) (CA INDEX NAME) C16 H16 N2 O6 S2 . Na CN

MF

LC STN Files: CA, CAPLUS

CRN (50674-67-4)

$$\langle s \rangle$$

$$\begin{array}{c} \text{D1-CH}_2-\text{C-NH} \\ \text{O} \\ \end{array}$$

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 40 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29

86005-06-3 REGISTRY RN

Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino], monolithium salt, (6R-cis)- (9CI) (CA INDEX NAME) CN

STEREOSEARCH

C16 H17 N3 O7 S2 . Li MF

STN Files: CA, CAPLUS, IMSPATENTS, USPATFULL LC

(35607-66-0) CRN

Absolute stereochemistry.

● Li

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 41 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

85643-59-0 REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-, (6R-trans)-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

Acetic acid, trifluoro-, compd. with (6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid (1:1)

FS STEREOSEARCH

MF C17 H17 N3 O6 S2 . C2 H F3 O2

STN Files: CA, CAPLUS

CM

21593-23-7 CRN CMF C17 H17 N3 O6 S2

Absolute stereochemistry.

CM

CRN 76-05-1 C2 H F3 02 CMF

CO<sub>2</sub>H

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 42 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN **85643-58-9** REGISTRY Entered STN: 16 Nov 1984 L29

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-, monohydrochloride, (6R-trans)- (9CI) (CA INDEX NAME) CN

STEREOSEARCH

MF C17 H17 N3 O6 S2 . C1 H

STN Files: CA, CAPLUS (21593-23-7) LC

CRN

Absolute stereochemistry.

HC1

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 43 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

85293-56-7 REGISTRY
Entered STN: 16 Nov 1984
5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-,
(6R-trans)-, sulfite (1:1) (9CI) (CA INDEX NAME)
STEREOSEARCH ED CN FS MF C17 H17 N3 O6 S2 . H2 O3 S STN Files: CA, CAPLUS CRN 21593-23-7 CMF C17 H17 N3 06 S2

Absolute stereochemistry.

CM

7782-99-2 CMF H2 03 S

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 44 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 82136-28-5 REGISTRY
ED Entered STN: 16 Nov 1984
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino], hydrate, (6R-cis)- (9CI) (CA INDEX NAME)
FS STEREOSEARCH

MF C16 H17 N3 O7 S2 . x H2 O

CRN (35607-66-0)

Absolute stereochemistry.

●x H<sub>2</sub>0

```
L29 ANSWER 45 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
      80823-78-5 REGISTRY
Entered STN: 16 Nov 1984
ED
      L-Lysine, mono[(6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[[(4-
      pyridinylthio)acetyl]amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-
      carboxylate] (9CI) (CA INDEX NAME)
- OTHER CA INDEX NAMES:
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-,
      (6R-trans)-, compd. with L-lysine (1:1) (9CI) STEREOSEARCH
FS
      C17 H17 N3 O6 S2 . C6 H14 N2 O2
MF
LC
      STN Files: CA, CAPLUS
      CM
      CRN
            21593-23-7
      CMF
            C17 H17 N3 06 S2
```

Absolute stereochemistry.

CM 2

CRN 56-87-1 CMF C6 H14 N2 O2

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 46 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 80813-08-7 REGISTRY Entered STN: 16 Nov 1984 L29 RN ED L-Cysteine, N-acetyl-, (6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate (salt) (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-, (6R-trans)-, compd. with N-acetyl-L-cysteine (1:1) (9CI) OTHER NAMES: CN Cephapirin acetylcysteinate FS STEREOSEARCH C17 H17 N3 O6 S2 . C5 H9 N O3 S STN Files: CA, CAPLUS, USPATFULL CM 1 21593-23-7 CRN CMF C17 H17 N3 06 S2

Absolute stereochemistry.

2 CM

616-91-1 CRN C5 H9 N O3 S CMF

Absolute stereochemistry.

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 47 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

80813-07-6 REGISTRY Entered STN: 16 Nov 1984 ED

L-Cysteine, N-acetyl-,  $[6R-[6\alpha,7\beta(R*)]]-7-[[amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-5-thia-1-azabicyclo[4.2.0]oct-2$ ene-2-carboxylate (salt) (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo- $[6R-[6\alpha, 7\beta(R*)]]$ -, compd. with N-acetyl-L-cysteine (1:1) (9CI) OTHER NAMES:

CN Cefadroxil N-acetylcysteinate

FS STEREOSEARCH

MF C16 H17 N3 O5 S . C5 H9 N O3 S

STN Files: CA, CAPLUS, IMSPATENTS, USPATFULL

CM

CRN 50370-12-2 C16 H17 N3 O5 S CMF

Absolute stereochemistry.

CM 2

616-91-1 CRN C5 H9 N 03 S CMF

Absolute stereochemistry.

# 2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 48 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 80525-84-4 REGISTRY L29

RN

Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-CN calcium salt (2:1), (6R)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

C16 H17 N3 O7 S2 . 1/2 Ca MF

LC STN Files: CA, CAPLUS, IMSPATENTS, TOXCENTER, USPATFULL

(40126-15-6) CRN

Absolute stereochemistry.

#### ●1/2 Ca

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 49 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**79226-63-4** REGISTRY Entered STN: 16 Nov 1984 ED

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-5-thiazolyl)(methoxyimino)acetyl]amino]-8-oxo-,  $[6R-[6\alpha,7\beta(Z)]]-(9CI)$  (CA INDEX NAME)

STEREOSEARCH FS

MF C13 H13 N5 O5 S2

LC STN Files: CA, CAPLUS, TOXCENTER

Absolute stereochemistry. Double bond geometry as shown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 50 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 75859-41-5 REGISTRY ED Entered STN: 16 Nov 1984

CN 5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-, monosodium salt,  $[6R-(6\alpha,7\beta)]-$  (9CI) (CA INDEX NAME)

STEREOSEARCH FS

C13 H13 N5 O5 S2 . Na MF

LC STN Files: CA, CAPLUS

(770660-21-4)CRN

Absolute stereochemistry. Double bond geometry unknown.

Na

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 51 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 75360-38-2 REGISTRY ED Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-,
hydrochloride, [6R-[6α, 7β(Z)]]- (9CI) (CA INDEX NAME) CN

FS STEREOSEARCH

ΜF C13 H13 N5 O5 S2 . x C1 H

CA, CAPLUS, IMSPATENTS, TOXCENTER LC STN Files:

CRN (68401-81-0)

●x HC1

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 52 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**75275-72-8** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-, monohydrochloride,  $[6R-[6\alpha, 7\beta(E)]]-(9CI)$  (CA INDEX NAME)

FS STEREOSEARCH

C13 H13 N5 O5 S2 . C1 H MF

STN Files: CA, CAPLUS, TOXCENTER (68403-31-6) LC

Absolute stereochemistry. Double bond geometry as shown.

HC1

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 53 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**74708-63-7** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino)acetyl]amino]-8-oxo-, (6R-trans)-(9CI) (CA INDEX NAME)

STEREOSEARCH FS

MF C13 H13 N5 O5 S2

CI COM

STN Files: BEILSTEIN\*, CA, CAPLUS, CASREACT, TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

Absolute stereochemistry. Double bond geometry unknown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1907 TO DATE) 5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 54 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

74708-62-6 REGISTRY Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-, monosodium salt, (6R-trans)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C13 H13 N5 O5 S2 . Na

LC STN Files: BEILSTEIN\*, CA, CAPLUS, CASREACT, TOXCENTER (\*File contains numerically searchable property data) CRN (74708-63-7)

Absolute stereochemistry. Double bond geometry unknown.

Na

3 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 55 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

74660-11-0 REGISTRY RN

Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[2-(aminooxy)-2-oxoethyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-cis)- (9CI) (CA INDEX NAME)

FS STEREOSEARCH C16 H17 N3 O7 S2 MF

STN Files: CA, CAPLUS

Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 56 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 73356-25-9 REGISTRY RN

Entered STN: 16 Nov 1984 ED

CN

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-,
(6R-trans)-, sulfate (salt) (9CI) (CA INDEX NAME)

STEREOSEARCH

MF C16 H17 N3 O7 S2 . x H2 O4 S

CA, CAPLUS STN Files:

> CM 1

73356-24-8 C16 H17 N3 O7 S2

Absolute stereochemistry.

CM

CRN 7664-93-9 CMF H2 04 S

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 57 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 73356-24-8 REGISTRY ED Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-

(6R-trans) - (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C16 H17 N3 O7 S2

CI COM

STN Files: BEILSTEIN\*

(\*File contains numerically searchable property data)

Absolute stereochemistry.

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

L29 ANSWER 58 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

71420-79-6 REGISTRY RN

Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-hydroxyphenylacetyl]amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-5-yl]thio]methyl]-, monosodium salt, (6R, 7R)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[(hydroxyphenylacetyl)amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-5-yl]thio]methyl]-, monosodium salt, [6R-[6\alpha, 7\beta(R\*)]]-

OTHER NAMES:

CN Cefodie

CN Monocidur

CN Mopnocid

CN Praticef

FS STEREOSEARCH

MF C18 H18 N6 O8 S3 . Na

CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, PHAR, STN Files: PROUSDDR, TOXCENTER, USAN, USPATFULL (61270-58-4)

CRN

Absolute stereochemistry.

Na

# \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

7 REFERENCES IN FILE CA (1907 TO DATE) 7 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 59 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

70356-03-5 REGISTRY ED Entered STN: 16 Nov 1984 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, monohydrate, (6R, 7R)-(9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[(aminophenylacetyl)amino]-3-chloro-8-oxo-, monohydrate,  $[6R-[6\alpha,7\beta(R*)]]-$ OTHER NAMES: Cefaclor monohydrate Cephaclor monohydrate FS STEREOSEARCH DR 126269-39-4 C15 H14 C1 N3 O4 S . H2 O MF STN Files: ANABSTR, BIOSIS, CA, CAPLUS, CHEMCATS, EMBASE, IMSPATENTS, IPA, MEDLINE, MRCK\*, PHAR, PROUSDDR, RTECS\*, SYNTHLINE, USAN, USPAT2, (\*File contains numerically searchable property data) (53994-73-3)

Absolute stereochemistry.

$$\begin{array}{c|c} & & & & \\ & & & \\ H_2N & & & \\ & & & \\ Ph & & \\ \end{array}$$

● H20

15 REFERENCES IN FILE CA (1907 TO DATE) 15 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 60 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN 69088-92-2 REGISTRY

ED Entered STN: 16 Nov 1984

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[(hydroxyphenylacetyl)amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-5-yl]thio]methyl]-, [6R-(6α, 7β)]- (9CI) (CA INDEX NAME)

FS STEREOSEARCH

MF C18 H18 N6 08 S3

CI COM

LC STN Files: BEILSTEIN\*, CA, CAPLUS, IMSPATENTS
(\*File contains numerically searchable property data)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE)

# 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

Absolute stereochemistry.

Double bond geometry as shown.

# ●1/2 Mg

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

# ●1/2 Ca

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
L29 ANSWER 63 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
     68401-82-1 REGISTRY Entered STN: 16 Nov 1984
RN
ED
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-,
CN
     monosodium salt, (6R, 7R) - (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
      7-[[(2-amino-4-thiazolyl)(methoxyimino)acetyl]amino]-8-oxo-, monosodium
salt, [6R-[6α, 7β(Z)]]-
OTHER NAMES:
CN
     Cefizox
CN
     Ceftix
     Ceftizon
     Ceftizoxime sodium
      Ceftizoxime sodium salt
CN
     Eposerin
CN
     FK 749
     FR 13479
CN
CN
     SKF 88373
     Sodium 7-[2-(2-aminothiazol-4-yl)-2-methoxyiminoacetamido]-3-cephem-4-
      carboxylate
      STEREOSEARCH
     C13 H13 N5 O5 S2 . Na
MF
CI
                     BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CAPLUS,
        CASREACT, CBNB, CHEMLIST, CIN, DIOGENES, EMBASE, IFICDB, IFIUDB, IMSPATENTS, IPA, MRCK*, MSDS-OHS, PHAR, PROMT, PS, RTECS*, TOXCENTER,
        USAN, USPATFULL
           (*File contains numerically searchable property data)
      (68401-81-0)
```

Na

95 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 95 REFERENCES IN FILE CAPLUS (1907 TO DATE) L29 ANSWER 64 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN **68401-81-0** REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2Z)-(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-, (6R, 7R)-(9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl] amino]-8-oxo-,  $[6R-[6\alpha, 7\beta(Z)]]-$ OTHER NAMES: Ceftisomin CN Ceftizoxime CN Epocelin FS STEREOSEARCH MF C13 H13 N5 O5 S2 CI COM ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN\*, STN Files: BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CHEMCATS, CIN, DDFU, DIOGENES, DRUGU, EMBASE, IFICDB, IFIUDB, IMSCOSEARCH, IMSPATENTS, IPA, MEDLINE, MRCK\*, NIOSHTIC, PHAR, PROMT, PS, RTECS\*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL, VETU (\*File contains numerically searchable property data) Other Sources: WHO

Absolute stereochemistry.

Double bond geometry as shown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1069 REFERENCES IN FILE CA (1907 TO DATE)
4 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
1070 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 65 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 68401-80-9 REGISTRY
ED Entered STN: 16 Nov 1984
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2-amino-4-thiazolyl) (methoxyimino) acetyl]amino]-8-oxo-,
monohydrochloride, [6R-[6α, 7β(Z)]]- (9CI) ·(CA INDEX NAME)
FS STEREOSEARCH
MF C13 H13 N5 05 S2 . C1 H
LC STN Files: CA, CAPLUS, IMSPATENTS, PS, TOXCENTER, USPATFULL
CRN (68401-81-0)

Absolute stereochemistry.

Double bond geometry as shown.

HC1

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 66 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 67022-10-0 REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[[(aminocarbonyl) oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl) amino]-, (6R-cis)-, mixt. with [R-(R\*, S\*)]-[1-[(2-amino-1-oxopropyl) amino]ethyl]phosphonic acid (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: Phosphonic acid, [1-[(2-amino-1-oxopropyl)amino]ethyl]-, [R-(R\*, S\*)]-, mixt. contg. (9CI) STEREOSEARCH C16 H17 N3 O7 S2 . C5 H13 N2 O4 P MF CIMXS CA, CAPLUS, IMSPATENTS STN Files: CM 1 60668-24-8 CRN C5 H13 N2 O4 P

Absolute stereochemistry.

CM

CRN 35607-66-0 C16 H17 N3 07 S2 CMF

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 67 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

66911-02-2 REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[[(aminocarbonyl)oxy]methyl]-6-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-(9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C16 H17 N3 07 S2

CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, USPATFULL STN Files:

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 68 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29

66592-87-8 REGISTRY RN

Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, monohydrate, (6R, 7R) - (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, monohydrate,  $[6R-[6\alpha, 7\beta(R*)]]-$ 

OTHER NAMES:

Cefadroxil monohydrate

CN Duricef

FS STEREOSEARCH

C16 H17 N3 O5 S . H2 O MF

STN Files: ANABSTR, BEILSTEIN\*, BIOBUSINESS, BIOSIS, CA, CAPLUS, CBNB,

CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, DIOGENES, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MEDLINE, MRCK\*, PROMT, RTECS\*, TOXCENTER, USAN, USPAT2, USPATFULL

(\*File contains numerically searchable property data)

(50370-12-2)

Absolute stereochemistry.

● H20

19 REFERENCES IN FILE CA (1907 TO DATE)
19 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 69 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

66065-78-9 REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-,
(6R-trans)-, sulfate (1:1) (9CI) (CA INDEX NAME)
STEREOSEARCH

FS

C17 H17 N3 O6 S2 . H2 O4 S MF

LC STN Files: CA, CAPLUS

CM

CRN 21593-23-7 CMF C17 H17 N3 O6 S2

Absolute stereochemistry.

CM

CRN 7664-93-9 CMF H2 04 S

3 REFERENCES IN FILE CA (1907 TO DATE)
3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

Qazi 10/642194 Page 43

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ANSWER 70 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
L29
        64544-07-6 REGISTRY
RN
        Entered STN: 16 Nov 1984
        5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-[[(2Z)-2-furanyl(methoxyimino)acetyl]amin
o]-8-oxo-, 1-(acetyloxy)ethyl ester, (6R, 7R)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
        5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-[[2-furanyl(methoxyimino)acetyl]amino]-8-
oxo-, 1-(acetyloxy)ethyl ester, [6R-[6α, 7β(Z)]]-
OTHER NAMES:
CN
        CCI 15641
CN
        Cefazine
CN
        Ceftin
CN
        Cefurax
CN
        Cefuroxime 1-acetoxyethyl ester
CN
        Cefuroxime axetil
CN
        Elobact
CN
        Oraxim
CN
CN
        Zinat
         Zinnat
        STEREOSEARCH
MF
        C20 H22 N4 010 S
CI
           IN Files: ADISNEWS, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CIN, DDFU, DIOGENES, DRUGU, EMBASE, IMSCOSEARCH, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, PHAR, PROMT, PROUSDDR, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL
        STN Files:
                 (*File contains numerically searchable property data)
```

Absolute stereochemistry.

Double bond geometry as shown.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

6 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
408 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 71 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 63824-86-2 REGISTRY
ED Entered STN: 16 Nov 1984
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino], monosodium salt, (6S-cis)- (9CI) (CA INDEX NAME)
FS STEREOSEARCH

407 REFERENCES IN FILE CA (1907 TO DATE)

MF C16 H17 N3 O7 S2 . Na

LC STN Files: BEILSTEIN\*, CA, CAPLUS, IMSPATENTS
(\*File contains numerically searchable property data)
CRN (56082-95-2)

Absolute stereochemistry.

Na

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 72 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 61270-78-8 REGISTRY RN Entered STN: 16 Nov 1984 ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[(2R)-hydroxyphenylacetyl]amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-5-yl]thio]methyl]-, disodium salt, (6R, 7R)- (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[(hydroxyphenylacetyl)amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-5yl]thio]methyl]-, disodium salt,  $[6R-[6\alpha,7\beta(R*)]]$ -OTHER NAMES: CN Cefonicid disodium salt CN Cefonicid sodium CN Disodium cefonicid CN SKF 75073 STEREOSEARCH FS C18 H18 N6 O8 S3 . 2 Na MF STN Files: ANABSTR, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CHEMCATS, CIN, DIOGENES, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MRCK\*, PHAR, PROMT, PROUSDDR, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL (\*File contains numerically searchable property data) CRN (61270-58-4)

Absolute stereochemistry.

#### ●2 Na

39 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
39 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 73 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN 61270-58-4 REGISTRY

ED Entered STN: 16 Nov 1984

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,

```
7-[[(2R)-hydroxyphenylacetyl]amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-
       5-yl]thio]methyl]-, (6R, 7R)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
       7-[(hydroxyphenylacetyl)amino]-8-oxo-3-[[[1-(sulfomethyl)-1H-tetrazol-5-
       yl]thio]methyl]-, [6R-[6\alpha, 7\beta(R*)]]-
OTHER NAMES:
CN
      Cefonicid
FS
      STEREOSEARCH
DR
      162143-29-5
      C18 H18 N6 O8 S3
MF
CI
      COM
         IN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CBNB, CIN, DDFU, DIOGENES, DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MEDLINE, MRCK*, PROMT, PROUSDDR, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL
      STN Files:
             (*File contains numerically searchable property data)
       Other Sources:
                              WHO
```

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

```
199 REFERENCES IN FILE CA (1907 TO DATE)
  4 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
200 REFERENCES IN FILE CAPLUS (1907 TO DATE)
```

- ANSWER 74 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN  $58648 \hbox{-} 57 \hbox{-} 0$  REGISTRY L29
- RN
- Entered STN: 16 Nov 1984 ED
- 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
  7-[(hydroxyphenylacetyl)amino]-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, monolithium salt,  $[6R-[6\alpha, 7\beta(R*)]]$ - (9CI) (CA INDEX NAME)

OTHER NAMES:

- CN Cefamandole lithium
- CN Cefamandole lithium salt
- FS STEREOSEARCH
- C18 H18 N6 O5 S2 . Li MF
- CI COM
- N Files: BIOSIS, CA, CAPLUS, CHEMCATS, CSCHEM, IFICDB, IFIPAT, IFIUDB, IMPATENTS, TOXCENTER, USPATFULL LC STN Files:

CRN (34444-01-4)

Absolute stereochemistry.

● Li

33 REFERENCES IN FILE CA (1907 TO DATE)
33 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 75 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**57320-88-4** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-7-(acetyl-2-thienylamino)-8-oxo-, sodium salt,

(6R-trans) - (9CI) (CA INDEX NAME)

STEREOSEARCH FS

МF C16 H16 N2 O6 S2 . Na

LC STN Files: CA, CAPLUS, TOXCENTER

(758628-65-8) CRN

Absolute stereochemistry.

Na

- 3 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)
- ANSWER 76 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 56540-85-3 REGISTRY Entered STN: 16 Nov 1984 L29

RN

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-, monosodium salt, (6R)- (9CI) (CA INDEX NAME)

STEREOSEARCH FS

MF C16 H17 N3 O7 S2 . Na STN Files: BEILSTEIN\*

LC

(\*File contains numerically searchable property data)

(40126-15-6)

Absolute stereochemistry.

Na

```
L29 ANSWER 77 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 56082-95-2 REGISTRY
ED Entered STN: 16 Nov 1984
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-,
(6S-cis)- (9CI) (CA INDEX NAME)
FS STEREOSEARCH
MF C16 H17 N3 07 S2
CI COM
LC STN Files: BEILSTEIN*, CA, CAPLUS, IMSPATENTS
```

(\*File contains numerically searchable property data)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
L29 ANSWER 78 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
      55944-32-6 REGISTRY
Entered STN: 16 Nov 1984
RN
ED
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-tcompd. with N,N-dimethylacetamide (1:1) (9CI) (CA INDEX NAME)
                                                                                (6R-trans)-,
OTHER CA INDEX NAMES:
      Acetamide, N, N-dimethyl-, (6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[(2-
      thienylacetyl)amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate
      (9CI)
      STEREOSEARCH
      C16 H16 N2 O6 S2 . C4 H9 N O
      STN Files: CA, CAPLUS, TOXCENTER
      CM
      CRN 153-61-7
      CMF C16 H16 N2 O6 S2
```

CM 2

CRN 127-19-5 CMF C4 H9 N O

Me Me— N— Ac

> 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

Absolute stereochemistry.

(746546 - 33 - 8)

Na

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 80 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 54544-02-4 REGISTRY
ED Entered STN: 16 Nov 1984
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
7-[[[(carboxymethyl)thio]acetyl]amino]-8-oxo-3-(3-pyridinylmethyl)-,
(6R-trans)-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN Acetic acid, trifluoro-, compd. with (6R-trans)-7[[[(carboxymethyl)thio]acetyl]amino]-8-oxo-3-(3-pyridinylmethyl)-5-thia-1azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid (1:1)
FS STEREOSEARCH
MF C17 H17 N3 06 S2 . C2 H F3 02
LC STN Files: BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB

(\*File contains numerically searchable property data)

CM

54544-01-3 CRN C17 H17 N3 06 S2

Absolute stereochemistry.

CM 2

CRN 76-05-1 CMF C2 H F3 02

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 81 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**54544-01-3** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[[(carboxymethyl)thio]acetyl]amino]-8-oxo-3-(3-pyridinylmethyl)-, (6R-trans) - (9CI) (CA INDEX NAME)

STEREOSEARCH FS

MF C17 H17 N3 O6 S2

CI COM

STN Files: BEILSTEIN\*

(\*File contains numerically searchable property data)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

ANSWER 82 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29

53994-73-3 REGISTRY RN

Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-aminophenylacetyl]amino]-3-chloro-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[(aminophenylacetyl)amino]-3-chloro-8-oxo-, [6R-[6α, 7β(R\*)]]-

```
OTHER NAMES:
        7-(D-2-Amino-2-phenylacetamido)-3-chloro-3-cephem-4-carboxylic acid
       Alfacet
        Ceclor
       Cefaclor
CN
        Cephaclor
       Kefral
       Lilly 99638
CN
       Panoral
CN
        S 6472
FS
        STEREOSEARCH
        142975-47-1
DR
MF
        C15 H14 C1 N3 O4 S
CI
        COM
           IN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM, DDFU, DIOGENES, DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSCOSEARCH, IMSPATENTS, IPA, MEDLINE, MSDS-OHS, PHAR, PROMT, PROUSDDR, PS, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL,
        STN Files:
           VETU
               (*File contains numerically searchable property data)
        Other Sources: EINECS**, WHO
               (**Enter CHEMLIST File for up-to-date regulatory information)
```

Absolute stereochemistry.

$$\begin{array}{c|c} & & & & & \\ & & & & \\ & & & & \\ H_2N & & & & \\ P_h & & & & \\ \end{array}$$

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**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
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```
20 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
1483 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 83 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 53950-16-6 REGISTRY
ED Entered STN: 16 Nov 1984
CN L-Arginine, mono[(6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate] (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-,
(6R-trans)-, compd. with L-arginine (1:1) (9CI)
FS STEREOSEARCH
MF C17 H17 N3 06 S2 . C6 H14 N4 O2
LC STN Files: CA, CAPLUS, TOXCENTER

CM 1

CRN 21593-23-7
```

1479 REFERENCES IN FILE CA (1907 TO DATE)

Absolute stereochemistry.

C17 H17 N3 06 S2

CMF

CM

CRN 74-79-3 CMF C6 H14 N4 O2

Absolute stereochemistry.

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 84 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 52128-54-8 REGISTRY

Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino], monosodium salt, cis- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]monosodium salt,  $cis^-(\pm)^-$ 

STEREOSEARCH FS

MF

C16 H17 N3 O7 S2 . Na STN Files: BEILSTEIN\*, CA, CAPLUS, CHEMCATS, IMSPATENTS, TOXCENTER LC (\*File contains numerically searchable property data) CRN (51014-44-9)

Relative stereochemistry.

Na

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 85 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

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51829-86-8 REGISTRY
     Entered STN: 16 Nov 1984
ED
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-
CN
      monosodium salt, (6R, 7R) - (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
     3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-
       monosodium salt, (6R-trans)-
OTHER NAMES:
     Sodium cefoxitin
CN
     STEREOSEARCH
     C16 H17 N3 O7 S2 . Na
STN Files: AGRICOLA, BEILSTEIN*, BIOBUSINESS, BIOSIS, CA, CAPLUS,
MF
LC
        IMSPATENTS, TOXCENTER, USPATFULL
          (*File contains numerically searchable property data)
CRN
    (73356-24-8)
```

Absolute stereochemistry.

Na

39 REFERENCES IN FILE CA (1907 TO DATE) 39 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
L29 ANSWER 86 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
      51596-58-8 REGISTRY
RN
      Entered STN: 16 Nov 1984
ED
      L-Arginine, mono[(6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate]
       (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-trans)-,
compd. with L-arginine (1:1) (9CI)
OTHER NAMES:
      Cephalothin arginine salt
CN
      STEREOSEARCH
      C16 H16 N2 O6 S2 . C6 H14 N4 O2
MF
                      BEILSTEIN*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER,
LC
      STN Files:
         USPATFULL
            (*File contains numerically searchable property data)
      CM
            1
      CRN 153-61-7
      CMF
            C16 H16 N2 O6 S2
```

CM

CRN 74-79-3 CMF C6 H14 N4 O2

Absolute stereochemistry.

3 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 87 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**51596-52-2** REGISTRY Entered STN: 16 Nov 1984 ED

L-Lysine, mono[(6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[(2thienylacetyl)amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate] (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-trans)-, compd. with L-lysine (1:1) (9CI)

OTHER NAMES:

Lysine 3-(acetoxymethyl)-7-(2-thienylacetamido)- $\Delta$ 3-cephem-4-CN carboxylate

STEREOSEARCH

MF

C16 H16 N2 O6 S2 . C6 H14 N2 O2 STN Files: BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER, LC USPATFULL (\*File contains numerically searchable property data)

CM 1

CRN 153-61-7 CMF C16 H16 N2 O6 S2

Absolute stereochemistry.

2 CM

CRN 56-87-1 CMF C6 H14 N2 02 Absolute stereochemistry.

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 88 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**51266-76-3** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-trans)-,
compd. with gentamicin (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

Gentamicin, compd. with (6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[(2thienylacetyl)amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid (9CI)

OTHER NAMES:

CN Gentamicin cephalothinate

FS STEREOSEARCH

C16 H16 N2 O6 S2 . x Unspecified STN Files: CA, CAPLUS, TOXCENTER MF LC

> CM 1

1403-66-3 CRN CMF Unspecified

CCI MAN

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

CM 2

CRN 153-61-7

CMF C16 H16 N2 O6 S2

Absolute stereochemistry.

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 89 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

51266-75-2 REGISTRY Entered STN: 16 Nov 1984 ED

Polymyxin, compd. with (6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[(2thienylacetyl)amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-trans)-, compd. with polymyxin (9CI)

OTHER NAMES:

Polymyxin cephalothinate CN

FS STEREOSEARCH

MF C16 H16 N2 O6 S2 . x Unspecified

CA, CAPLUS STN Files:

CM 1

CRN 1406-11-7 CMF Unspecified

CCI MAN

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

CM

CRN 153-61-7

CMF C16 H16 N2 06 S2

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 90 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**51098-29-4** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, trans- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, trans-(±)-

OTHER NAMES:

(±)-Cephalothin CN STEREOSEARCH

FS MF C16 H16 N2 O6 S2

CI COM

BEILSTEIN\*, BIOSIS, CA, CAPLUS, CASREACT, CHEMCATS, IFICDB, IFIPAT, IFIUDB, SYNTHLINE, USPATFULL

(\*File contains numerically searchable property data)

Relative stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1907 TO DATE) 5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 91 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**51014-52-9** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, monosodium salt, trans- (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES:

CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,

Page 56

3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, monosodium salt, trans-(±)-STEREOSEARCH

FS

C16 H16 N2 O6 S2 . Na MF

BEILSTEIN\*, CA, CAPLUS, CASREACT, CHEMCATS, IFICDB, IFIPAT, LC STN Files: IFIUDB, USPATFULL

(\*File contains numerically searchable property data)

(51098-29-4)

Relative stereochemistry.

#### Na

4 REFERENCES IN FILE CA (1907 TO DATE) 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 92 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

51014-44-9 REGISTRY RN

Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]cis- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]cis-(±)-

FS STEREOSEARCH MF C16 H17 N3 O7 S2

CI COM

BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, STN Files: TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

Relative stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1907 TO DATE) 5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 93 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29

RN

50674-67-4 REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(thienylacetyl)amino]-, (6R-trans)- (9CI) CN

(CA INDEX NAME)
MF C16 H16 N2 O6 S2

CI IDS, COM

LC STN Files: CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER, USPATFULL

$$\langle \hat{} \rangle$$

$$\begin{array}{c} \text{D1-CH}_2-\text{C-NH} \\ \vdots \\ \text{CO}_2\text{H} \end{array}$$

3 REFERENCES IN FILE CA (1907 TO DATE)
3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
3 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L29
      ANSWER 94 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
      50370-12-2 REGISTRY
RN
      Entered STN: 16 Nov 1984
ED
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
CN
      7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, (6R, 7R)-1
       (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
       7-[[amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-,
       [6R-[6\alpha, 7\beta(R*)]]-
OTHER NAMES:
      Bidocef
CN
CN
      BL-S 578
CN
      Cefadroxil
      Cephadroxi1
CN
      D-Cefadroxil
CN
      Oracefal
      Roxil
CN
      S 578
CN
      Ultracef
FS
      STEREOSEARCH
      C16 H17 N3 O5 S
CI
      COM
         IN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM, DDFU, DIOGENES, DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSCOSEARCH, IMSPATENTS, MEDLINE, PHAR, PROMT, PS, RTECS*, TOXCENTER, USAN, USPAT2, USPATFULL, VETU, VTB
      STN Files:
            (*File contains numerically searchable property data)
                            EINECS**, WHO
      Other Sources:
            (**Enter CHEMLIST File for up-to-date regulatory information)
```

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

Qazi 10/642194 Page 58

879 REFERENCES IN FILE CA (1907 TO DATE)
19 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
881 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 95 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29 **47489-80-5** REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-,  $[2R-(2\alpha, 6\alpha, 7\beta)]-(9CI)$  (CA INDEX NAME) STEREOSEARCH C16 H16 N2 O6 S2 MF COM CI BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER, LC STN Files: USPATFULL (\*File contains numerically searchable property data)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

4 REFERENCES IN FILE CA (1907 TO DATE) 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 96 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 42284-83-3 REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, monosodium salt, (6R, 7R) - (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[[amino(4-hydroxyphenyl)acetyl]amino]-3-methyl-8-oxo-, monosodium salt,  $[6R-[6\alpha, 7\beta(R*)]]-$ OTHER NAMES: p-Hydroxycephalexin CN Sodium cefadroxil STEREOSEARCH C16 H17 N3 O5 S . Na CA, CAPLUS, CHEMLIST, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, STN Files: TOXCENTER, USPATFULL

Na

10 REFERENCES IN FILE CA (1907 TO DATE) 10 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 97 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN **42231-00-5** REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-trans)-, homopolymer (9CI) (CA INDEX NAME) CN OTHER NAMES: Cephalothin polymer CN STEREOSEARCH FS 100296-27-3 DR MF (C16 H16 N2 06 S2)xCI **PMS** Polyamine, Polyester, Polyester formed **PCT** STN Files: ADISNEWS, AGRICOLA, CA, CAPLUS, DIOGENES, MEDLINE, PROMT CM 1 CRN 153-61-7

Absolute stereochemistry.

CMF

C16 H16 N2 O6 S2

2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

(\*File contains numerically searchable property data)

ANSWER 98 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 40126-15-6 REGISTRY RN Entered STN: 16 Nov 1984 ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-(6R) - (9CI) (CA INDEX NAME) STEREOSEARCH FS MF C16 H17 N3 O7 S2 CI COM BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, LC STN Files: TOXCENTER, USPATFULL

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

3 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

CN Ethanamine, N, N-diethyl-, (6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate

FS STEREOSEARCH

MF C16 H16 N2 O6 S2 . C6 H15 N

LC STN Files: CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER

CM 1

CRN 153-61-7 CMF C16 H16 N2 06 S2

Absolute stereochemistry.

CM 2

CRN 121-44-8 CMF C6 H15 N

Et | Et—N—Et

> 3 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 100 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN 35607-66-0 REGISTRY
ED Entered STN: 16 Nov 1984
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-

, (6R,7S)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:
CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,

```
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-
        (6R-cis)-
OTHER NAMES:
CN
      Cefoxitin
CN
      Cephoxitin
FS
      STEREOSEARCH
      39951-67-2
DR
MF
      C16 H17 N3 O7 S2
CI
      COM
        TN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAPLUS, CASREACT, CBNB, CHEMLIST, CIN,
      STN Files:
        CSCHEM, CSNB, DDFU, DIOGENES, DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB,
        IMSCOSEARCH, IMSPATENTS, IPA, MEDLINE, MRCK*, NAPRALERT, PROMT, PS, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL, VETU
           (*File contains numerically searchable property data)
      Other Sources: EINECS**, WHO
           (**Enter CHEMLIST File for up-to-date regulatory information)
```

Absolute stereochemistry.

```
**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
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Other Sources: EINECS\*\*, WHO

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3067 REFERENCES IN FILE CA (1907 TO DATE)
17 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
3075 REFERENCES IN FILE CAPILIS (1907 TO DATE)
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3075 REFERENCES IN FILE CAPLUS (1907 TO DATE)
      ANSWER 101 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
       34444-01-4 REGISTRY
      Entered STN: 16 Nov 1984
ED
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
       7-[[(2R)-hydroxyphenylacetyl]amino]-3-[[(1-methyl-1H-tetrazol-5-
       yl)thio]methyl]-8-oxo-, (6R, 7R)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
       7-[(hydroxyphenylacetyl)amino]-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, [6R-[6\alpha, 7\beta(R*)]]-
OTHER NAMES:
       3-(1-\texttt{Methyltetrazol}-5-\texttt{ylthiomethyl})-7-\texttt{D-mandelamido}-3-\texttt{cephem-4-carboxylic}
       acid
       Cefadole
       Cefamandol
CN
       Cefamandole
       Cephadole
       Cephamandole
CN
CN
       CMT
CN
       L-Cefamandole
       STEREOSEARCH
FS
MF
       C18 H18 N6 O5 S2
CI
       COM
         IN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CIN, CSNB, DDFU, DIOGENES, DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MEDLINE, MRCK*, NAPRALERT, PHAR, PROMT, PS, RTECS*, SYNTHLINE, TOXCENTER, USAN, USPATZ, USPATFULL, VETU
       STN Files:
             (*File contains numerically searchable property data)
```

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

Absolute stereochemistry.

```
Ph
HO R R S N N
CO<sub>2</sub>H Me
```

## \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

```
1637 REFERENCES IN FILE CA (1907 TO DATE)
9 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
```

```
1642 REFERENCES IN FILE CAPLUS (1907 TO DATE)
     ANSWER 102 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
L29
      33564-30-6 REGISTRY
Entered STN: 16 Nov 1984
RN
ED
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-
CN
        monosodium salt, (6R, 7S) - (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
      3-[[(aminocarbonyl)oxy]methyl]-7-methoxy-8-oxo-7-[(2-thienylacetyl)amino]-
         monosodium salt, (6R-cis)-
OTHER NAMES:
      Betacef
      Cefaxilin sodium
      Cefoxitin sodium
CN
CN
      Cefoxitin sodium salt
CN
      Cenomycin
CN
      Farmoxin
CN
      Mefoxin
CN
      Mefoxithin
      Mefoxitin
CN
      Merxin
CN
      MK 306
CN
      Monosodium cefoxitin
      STEREOSEARCH
DR
      41224-60-6
      C16 H17 N3 O7 S2 . Na
MF
CI
      COM
        IN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CAPLUS, CASREACT, CBNB, CHEMCATS, CHEMLIST, CIN, DIOGENES, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MRCK*, MSDS-OHS, PHAR, PIRA, PROMT, PS, RTECS*, TOXCENTER, USAN, USPAT2,
         USPATFULL
            (*File contains numerically searchable property data)
      Other Sources: EINECS**
            (**Enter CHEMLIST File for up-to-date regulatory information)
      (35607-66-0)
```

Na

148 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 148 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 103 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29 RN 30034-03-8 REGISTRY ED Entered STN: 16 Nov 1984 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 7-[[(2R)-hydroxyphenylacetyl]amino]-3-[[(1-methyl-1H-tetrazol-5-methylyl)thio]methyl]-8-oxo-, monosodium salt, (6R, 7R)- (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-mandelamido-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, monosodium salt, D- (8CI) CN 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 7-[(hydroxyphenylacetyl)amino]-3-[[(1-methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-, monosodium salt, [6R-[6 $\alpha$ , 7 $\beta$ (R\*)]]-OTHER NAMES: Cefamandole sodium

- CN
- Cefamandole sodium salt CN
- Sodium 7-D-mandelamido-3-(1-methyl-1H-tetrazol-5-ylthiomethyl)-3-cephem-4-CN carboxylate
- Sodium cefamandole
- Sodium  $[6R-[6\alpha, 7\beta(R*)]]-7-[(hydroxyphenylacetyl)amino]-3-[[(1-$ CN methyl-1H-tetrazol-5-yl)thio]methyl]-8-oxo-5-thia-1-azabicyclo[4.2.0]oct-2ene-2-carboxylate
- FS STEREOSEARCH
- MF C18 H18 N6 O5 S2 . Na
- CI COM
- STN Files: BEILSTEIN\*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CIN, DIOGENES, EMBASE, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, RTECS\*, TOXCENTER, USAN, USPATFULL LC (\*File contains numerically searchable property data)

EINECS\*\* Other Sources:

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

CRN (34444-01-4)

## 62 REFERENCES IN FILE CA (1907 TO DATE) 62 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
L29 ANSWER 104 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
RN
      24412-14-4 REGISTRY
ED
      Entered STN: 16 Nov 1984
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester),
monosilver(1+) salt (8CI) (CA INDEX NAME)
CN
OTHER NAMES:
      Silver cephalothin STEREOSEARCH
CN
FS
DR
      25047-35-2
      C16 H16 N2 O6 S2 . Ag
MF
LC
                       CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER
      STN Files:
CRN
      (153-61-7)
```

Absolute stereochemistry.

## ● Ag(I)

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

```
L29 ANSWER 105 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
      24356-60-3 REGISTRY Entered STN: 16 Nov 1984
ED
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-,
monosodium salt, (6R, 7R)- (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      5-Thia-I-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(4-pyridylthio)acetamido]-, acetate (ester),
       monosodium salt (8CI)
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-,
       monosodium salt, (6R-trans)-
OTHER NAMES:
       7-[\alpha-(4-Pyridylthio)acetamido] cephalosporanic acid sodium salt
CN
       Ambrocef
       BLP 1322
       Brisfirina
       Bristocef
CN
       Cef-Lak
       Cefa-Lak
       Cefadyl
CN
       Cefaloject
CN
       Cefapirin sodium
       Cefaprin sodium
CN
CN
       Cefatrexyl
CN
       Cephapirin sodium
       Cephatrexil
       NSC 179171
       Piricef
CN
       Sodium cefapirin
CN
       Sodium cephapirin
CN
       Today
       STERÉOSEARCH
FS
       C17 H17 N3 O6 S2 . Na
```

CI N Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN\*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CIN, CSCHEM, DIOGENES, EMBASE, HSDB\*, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MRCK\*, STN Files: PROMT, PS, RTECS\*, TOXCENTER, USAN, USPAT2, USPATFULL (\*File contains numerically searchable property data) EINECS\*\* Other Sources: (\*\*Enter CHEMLIST File for up-to-date regulatory information) CRN (21593-23-7)

Absolute stereochemistry.

Na

126 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 126 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 106 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29

RN

22739-71-5 REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester),
compd. with diethylamine (1:1) (8CI) (CA INDEX NAME) CN OTHER CA INDEX NAMES:

Diethylamine, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate acetate (ester) (8CI)

STEREOSEARCH

C16 H16 N2 O6 S2 . C4 H11 N MF

LC STN Files: CA, CAPLUS

> CM 1

CRN 153-61-7 C16 H16 N2 O6 S2 CMF

Absolute stereochemistry.

CM

109-89-7 C4 H11 N CMF

H3C-CH2-NH-CH2-CH3

# 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 107 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 22612-01-7 REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, CN 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester), compd. with 1-deoxy-1-(methylamino)glucitol (1:1) (8CI) (CA INDEX NAME)

OTHER CA INDEX NAMES: Glucitol, 1-deoxy-1-(methylamino)-, 3-(hydroxymethyl)-8-oxo-7-[2-(2thienyl)acetamido]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate 3-acetate (salt) (8CI)

STEREOSEARCH

MF C16 H16 N2 O6 S2 . C7 H17 N O5

LC STN Files: CA, CAPLUS

CM

CRN 6284-40-8 C7 H17 N 05 CMF

Absolute stereochemistry.

CRN 153-61-7 CMF C16 H16 N2 O6 S2

Absolute stereochemistry.

## 1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 108 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN **22611-99-0** REGISTRY Entered STN: 16 Nov 1984 RN

ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester), compd. with 2, 2'-iminodiethanol (1:1) (8CI) (CA INDEX NAME)

OTHER CA INDEX NAMES: Ethanol, 2, 2'-iminodi-, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate 3-acetate (salt) (8CI)

STEREOSEARCH

C16 H16 N2 O6 S2 . C4 H11 N O2 MF

STN Files: CA, CAPLUS

CM 1

CRN 153-61-7 CMF C16 H16 N2 06 S2

CM 2

CRN 111-42-2 CMF C4 H11 N O2

HO-CH2-CH2-NH-CH2-CH2-OH

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 109 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 22199-74-2 REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, compd. with 2-aminoethanol (1:1) (9CI) (CA INDEX NAME) (6R-trans)-OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester), compd. with 2-aminoethanol (1:1) (8CI) Ethanol, 2-amino-, (6R-trans)-3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-5-thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylate (salt) (9CI) OTHER NAMES: 7-(2-Thienylacetamido) cephalosporanic acid ethanolamine salt CN FS STEREOSEARCH DR 22611-98-9 C16 H16 N2 O6 S2 . C2 H7 N O MF STN Files: CA, CAPLUS CM

Absolute stereochemistry.

CRN 153-61-7

CMF C16 H16 N2 O6 S2

CM 2

CRN 141-43-5 CMF C2 H7 N 0

H2N-CH2-CH2-OH

1 REFERENCES IN FILE CA (1907 TO DATE)

#### 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 110 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 21593-24-8 REGISTRY Entered STN: 16 Nov 1984 ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(3-pyridinylthio)acetyl]amino]-, (6R-trans) - (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(3-pyridylthio)acetamido]-, acetate (ester) STEREOSEARCH FS MF C17 H17 N3 O6 S2 LC STN Files: BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER (\*File contains numerically searchable property data)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

```
4 REFERENCES IN FILE CA (1907 TO DATE)
                      4 REFERENCES IN FILE CAPLUS (1907 TO DATE)
      ANSWER 111 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
      21593-23-7 REGISTRY
Entered STN: 16 Nov 1984
RN
ED
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-, (6R, 7R)-
       (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
       3-(hydroxymethyl)-8-oxo-7-[2-(4-pyridylthio)acetamido]-, acetate (ester)
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(4-pyridinylthio)acetyl]amino]-,
CN
       (6R-trans)-
OTHER NAMES:
       7-[2-(4-Pyridylthio)acetamido]cephalosporanic acid
CN
       Cefapirin
CN
       Cefaprin
CN
       Cephapirin
CN
       Cephapirine
CN
       Cephaprin
       STEREOSEARCH
DR
       25279-35-0
       C17 H17 N3 O6 S2
MF
CI
       COM
          IN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CHEMLIST, DDFU, DIOGENES, DRUGU, EMBASE, HSDB*, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MEDLINE, NAPRALERT, NIOSHTIC, PROMT, PS, RTECS*, TOXCENTER, USAN, USPAT2, USPATFULL, VETU
```

(\*File contains numerically searchable property data)

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

Absolute stereochemistry.

Other Sources: EINECS\*\*, WHO

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

557 REFERENCES IN FILE CA (1907 TO DATE)
8 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
561 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L29 ANSWER 112 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN 21593-22-6 REGISTRY Entered STN: 16 Nov 1984 RN ED 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[[(2-pyridinylthio)acetyl]amino]-, (6R-trans)- (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(2-pyridylthio)acetamido]-, acetate (ester) (8CI) OTHER NAMES: CN NSC 270289 STEREOSEARCH FS MF C17 H17 N3 O6 S2 STN Files: BEILSTEIN\*, CA, CAPLUS, IFICDB, IFIPAT, IFIUDB, TOXCENTER LC

(\*File contains numerically searchable property data)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

5 REFERENCES IN FILE CA (1907 TO DATE) 5 REFERENCES IN FILE CAPLUS (1907 TO DATE)

ANSWER 113 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN L29 **3432-90-4** REGISTRY RN Entered STN: 16 Nov 1984
5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[(3-thienylacetyl)amino]-, (6R, 7R)- (9CI) ED (CA INDEX NAME) OTHER CA INDEX NAMES: 5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-(hydroxymethyl)-8-oxo-7-[2-(3-thienyl)acetamido]-, acetate (ester) (8CI)
5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[[2-(3-thienyl)acetyl]amino]-, (6R-trans)-CN STEREOSEARCH FS C16 H16 N2 O6 S2 MF LC STN Files: BEILSTEIN\*, CA, CAOLD, CAPLUS, TOXCENTER, USPATFULL (\*File contains numerically searchable property data)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

- 3 REFERENCES IN FILE CA (1907 TO DATE) 3 REFERENCES IN FILE CAPLUS (1907 TO DATE) 1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
- L29 ANSWER 114 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN RN 1051-68-9 REGISTRY ED Entered STN: 16 Nov 1984

5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, monosodium salt,  $[6R-(6\alpha,7\beta)]-(9CI)$  (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester), monosodium salt (8CI)

5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate, sodium salt (7CI)

STEREOSEARCH

C16 H16 N2 O6 S2 . Na MF

BEILSTEIN\*, CA, CAOLD, CAPLUS, IFICDB, IFIPAT, IFIUDB, LC STN Files: USPATFULL

(\*File contains numerically searchable property data)

(979 - 94 - 2)

Absolute stereochemistry.

Na

- 2 REFERENCES IN FILE CA (1907 TO DATE) 2 REFERENCES IN FILE CAPLUS (1907 TO DATE) 6 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

ANSWER 115 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN

RN

**979-95-3** REGISTRY Entered STN: 16 Nov 1984 ED

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, monopotassium salt, (6R-trans)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester), monopotassium salt (8CI)

```
OTHER NAMES:
CN Cephalothin potassium salt
CN Potassium 7-(2-thienylacetamido)cephalosporanate
FS STEREOSEARCH
MF C16 H16 N2 O6 S2 . K
LC STN Files: BEILSTEIN*, CA, CAOLD, CAPLUS, IFICDB, IFIPAT, IFIUDB,
TOXCENTER, USPATFULL
(*File contains numerically searchable property data)
CRN (153-61-7)
```

Absolute stereochemistry.

K

```
6 REFERENCES IN FILE CA (1907 TO DATE)
                 6 REFERENCES IN FILE CAPLUS (1907 TO DATE)
                 1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
     ANSWER 116 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
     979-94-2 REGISTRY
RN
     Entered STN: 16 Nov 1984
ED
     5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid,
     3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R, 7R)-(9CI)
      (CA INDEX NAME)
OTHER CA INDEX NAMES:
     5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid,
     3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester) (8CI) 5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid,
     3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (7CI)
     5-Thia-1-azabicyclo[4.2.0]oct-3-ene-2-carboxylic acid, 3-[(acctyl) methyl]-8-oxo-7-[(2-thienylacetyl) amino]-,
     [6R-(6\alpha, 7\beta)]
OTHER NAMES:
     3-Acetoxymethyl-7-thiopheneacetamido-2-cephem-4-carboxylic acid
     STEREOSEARCH
DR
     94094-06-1, 96265-55-3
     C16 H16 N2 O6 S2
MF
CI
     COM
                    BEILSTEIN*, CA, CAOLD, CAPLUS, CASREACT, IFICDB, IFIPAT,
     STN Files:
        IFIUDB, TOXCENTER, USPATFULL
          (*File contains numerically searchable property data)
```

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

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1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
      ANSWER 117 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
      153-61-7 REGISTRY
RN
ED
       Entered STN: 16 Nov 1984
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R, 7R)- (9CI)
CN
       (CA INDEX NAME)
OTHER CA INDEX NAMES:
      5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid, 3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester) (8CI)
       5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
       3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, (6R-trans)-
OTHER NAMES:
      3-(Acetoxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-5-thia-1-2-(2-thienyl)acetamido]-5-thia-1-2-(2-thienyl)acetamido]-5-thia-1-2-(2-thienyl)acetamido]
       azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid
       3-Acetoxymethyl-7-(2-thienylacetamido)-3-cephem-4-carboxylic acid
       7-(2-Thienylacetamido) cephalosporanic acid
CN
CN
      7-(Thiophene-2-acetamido) cephalosporin
CN
       7-[2-(2-Thienyl) acetylamido] cephalosporanic acid
CN
      Cefalotin
CN
       Cephalothin
CN
      Cephalotin
CN
       CT
FS
       STEREOSEARCH
DR
       2073-29-2
       C16 H16 N2 O6 S2
MF
CI
       COM
         N Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB, CHEMCATS,
LC
       STN Files:
          CHEMLIST, CIN, CSCHEM, CSNB, DDFU, DIOGENES, DRUGU, EMBASE, HSDB*,
         IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MEDLINE, MRCK*, NAPRALERT, NIOSHTIC, PROMT, PS, RTECS*, TOXCENTER, USAN, USPAT2, USPATFULL, VETU (*File contains numerically searchable property data)
```

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

16 REFERENCES IN FILE CA (1907 TO DATE)
16 REFERENCES IN FILE CAPLUS (1907 TO DATE)

Absolute stereochemistry.

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

EINECS\*\*, WHO

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4122 REFERENCES IN FILE CA (1907 TO DATE)
              37 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
            4128 REFERENCES IN FILE CAPLUS (1907 TO DATE)
              43 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
    ANSWER 118 OF 118 REGISTRY COPYRIGHT 2005 ACS on STN
     58-71-9 REGISTRY
RN
     Entered STN: 16 Nov 1984
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
     3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, monosodium salt,
(6R, 7R) - (9CI)
OTHER CA INDEX NAMES:
                     (CA INDEX NAME)
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
     3-(hydroxymethyl)-8-oxo-7-[2-(2-thienyl)acetamido]-, acetate (ester),
     monosodium salt (8CI)
     5-Thia-1-azabicyclo[4.2.0]oct-2-ene-2-carboxylic acid,
```

```
3-[(acetyloxy)methyl]-8-oxo-7-[(2-thienylacetyl)amino]-, monosodium salt,
     (6R-trans)-
OTHER NAMES:
CN
     38253
CN
     Averon 1
CN
     Cefalothin sodium
CN
     Cefalothine sodium
CN
     Cemastin
CN
CN
     Cephalothin sodium
     Cephalothin sodium salt
CN
     Cephation
CN
     Ceporacin
CN
     Cepovenin
CN
     Coaxin
CN
     Keflin
CN
     Lilly 38253
CN
     Lospoven
CN
     Microtin
CN
     Seffin
     Sodium (thienylacetamido)cephalosporanate
CN
     Sodium 3-acetoxymethyl-7β-(2-thienylacetamido)ceph-3-em-4-carboxylate
CN
     Sodium 7-(2-thienylacetamido)-3-acetoxymethyl-3-cephem-4-carboxylate
CN
     Sodium 7-[2-(2-thienyl)acetamido]cephalosporanate
CN
CN
     Sodium cefalotin
CN
     Sodium cephalothin
CN
     Sodium cephalotin
CN
     Synclotin
CN
     Toricelocin
FS
     STEREOSEARCH
     11048-05-8, 63-60-5, 53493-25-7, 115699-56-4, 297760-78-2
DR
     C16 H16 N2 O6 S2 . Na
MF
CI
     COM
     STN Files: ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS,
LC
       BIOTECHNO, CA, CABA, CAOLD, CAPLUS, CASREACT, CHEMCATS, CHEMLIST, CIN,
       CSCHEM, DIOGENES, EMBASE, HSDB*, IFICDB, IFIPAT, IFIUDB, IMSPATENTS, IPA, MRCK*, NAPRALERT, PROMT, PS, RTECS*, TOXCENTER, USAN, USPAT2,
       USPATFULL
          (*File contains numerically searchable property data)
     Other Sources: EINECS**
          (**Enter CHEMLIST File for up-to-date regulatory information)
CRN
     (153-61-7)
Absolute stereochemistry.
```

Na

588 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
588 REFERENCES IN FILE CAPLUS (1907 TO DATE)
14 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> d ide 132 tot

L32 ANSWER 1 OF 2 REGISTRY COPYRIGHT 2005 ACS on STN RN 11138-66-2 REGISTRY ED Entered STN: 16 Nov 1984 CN Xanthan gum (9CI) (CA INDEX NAME)

```
OTHER NAMES:
       Actigum CX 9
       ADM 40
       B 1459
       Biopolymer 9702
CN
       Biopolymer XB 23
CN
       Biozan R
CN
       Bisfect XA 200
CN
       Bistop
CN
       Chemicogel
CN
       Duovis
CN
       E 415
CN
       Echogum
CN
       Echogum F
CN
       Echogum GM
CN
       Echogum RD
CN
       Echogum SF
CN
       Echogum T
CN
       Ekogum ketorol
CN
       Enorflo X
CN
       Flocon 1035
CN
       Flocon 4800
       Flocon 4800C
CN
CN
       Flodrill S
CN
       Galaxy XB
CN
       Gumixan K
CN
       Gums, xanthomonas
CN
       Idvis
       Inagel V 10
Inagel V 10K
CN
CN
CN
       Jungbunzlauer ST
CN
       K 5C151
CN
       K 9C57
CN
       Kelco BT
CN
       Kelflo
CN
       Keltrol
CN
       Keltrol BT
       Keltrol CG
CN
CN
       Keltrol F
CN
       Keltrol K 5C151
CN
       Keltrol RD
CN
       Keltrol SF
CN
       Keltrol T
       Keltrol TF
Keltrol TF 1000
CN
CN
CN
       Kelzan
       Kelzan 140X
CN
CN
       Kelzan AR
CN
       Kelzan ASX
       Kelzan ASXT
CN
       Kelzan D
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
       DISPLAY
       478314-71-5, 12673-42-6, 12771-06-1, 9088-32-8, 54511-23-8, 56592-13-3, 98112-77-7, 51811-95-1, 37189-49-4, 37279-85-9, 37332-19-7, 37383-52-1, 80450-59-5, 85568-76-9, 82600-55-3, 39393-27-6, 39444-54-7
       Unspecified
       PMS, COM, MAN
CI
       Manual registration, Polyester, Polyester formed
STN Files: AGRICOLA, ANABSTR, AQUIRE, BIOBUSINESS, BIOSIS, BIOTECHNO,
CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CHEMLIST, CIN,
CSCHEM, DDFU, DRUGU, EMBASE, ENCOMPLIT, ENCOMPLIT2, ENCOMPPAT,
ENCOMPPAT2, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*, MSDS-OHS,
NAPRALERT, NIOSHTIC, PIRA, PROMT, TOXCENTER, TULSA, USPAT2, USPATFULL,
             (*File contains numerically searchable property data)
       Other Sources: DSL**, EINECS**, TSCA**
              (**Enter CHEMLIST File for up-to-date regulatory information)
```

#### \*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

8792 REFERENCES IN FILE CA (1907 TO DATE)

295 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

8814 REFERENCES IN FILE CAPLUS (1907 TO DATE) L32 ANSWER 2 OF 2 REGISTRY COPYRIGHT 2005 ACS on STN 9000-30-0 REGISTRY RN Entered STN: 16 Nov 1984 Guar gum (9CI) (CA INDEX NAME) CN OTHER NAMES:  $\alpha\text{-}D\text{-}galacto\text{-}\beta\text{-}D\text{-}Mannan$ CN α-D-Galactopyrano-β-D-mannopyranan CN CN 1212A CN Avicel CE 15 CN Burtonite V 7E CN C 1000 CN C 1000 (gum) CN C 250 C 250 (gum) CN CN Celbond 7 Celca-Gum D 49D CN CN CP 3300 CN CSA-M 175 CN CSAA-M 80 CN Cyamopsis gum CN Dealca TP 1 CN Dealca TP 2 CN Decorpa CN Dycol 4500 CN E 412 CN **EGMB** Emcogum CSAA CN CN Emulgum 200 Emulgum 200S FFH 200 CN CN CN FG-HV Fine Gum G CN CN Fine Gum G 17 CN Frimulsion BM G 50 CN Galactasol Galactasol 20H5FI CN CN Galactasol 211 CN Galactasol 270 CN Galactasol 30M1F CN Galaxy 1083 CN Gendril Thik CN Gendriv 162 CN Guapack PF 20 CN Guapack PN CN Guar CN Guar 5200 Guar flour Guar HV 7000 CPS CN CN Guar Supercol U Fine Guaran CN Guarcel 302 CN Guarcol U 40 CN Guargel D 15 CN Gum cyamopsis CN ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for DISPLAY 9008-17-7, 9010-50-8, 9049-33-6, 9066-07-3, 53986-27-9, 57406-68-5, 57406-71-0, 63799-54-2, 85510-16-3 DR MF Unspecified PMS, COM, MAN CI Manual registration, Polyother, Polyother only STN Files: ADISNEWS, AGRICOLA, ANABSTR, AQUIRE, BIOBUSINESS, BIOSIS, PCT

BIOTECHNO, CA, CABA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM, CSNB, DDFU, DIOGENES, DRUGU, EMBASE, HSDB\*, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK\*, MSDS-OHS, NAPRALERT, NIOSHTIC, PHAR, PIRA, PROMT, RTECS\*, TOXCENTER, TULSA, USPAT2, USPATFULL, VTB

(\*File contains numerically searchable property data)

Other Sources: DSL\*\*, EINECS\*\*, TSCA\*\*

(\*\*Enter CHEMLIST File for up-to-date regulatory information)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*

\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

6618 REFERENCES IN FILE CA (1907 TO DATE)
822 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
6632 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d ide 140 tot

L40 ANSWER 1 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN
RN 827605-46-9 REGISTRY
ED Entered STN: 09 Feb 2005
CN 2-Propenoic acid, 2-methyl-, methyl ester, polymer with ethyl
2-propenoate, triblock (9CI) (CA INDEX NAME)
MF (C5 H8 02 . C5 H8 02) x
CI PMS
PCT Polyacrylic
SR CA

CM 1

CRN 140-88-5
CMF C5 H8 02

0 || EtO-C-CH=CH2

CM 2

CRN 80-62-6 CMF C5 H8 O2

H2C 0 || || Me=C=C=OMe

```
ANSWER 2 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN
L40
      736998-33-7 REGISTRY
Entered STN: 01 Sep 2004
RN
      2-Propenoic acid, 2-methyl-, methyl ester, polymer with ethyl 2-propenoate, diblock (9CI) (CA INDEX NAME) (C5 H8 02 . C5 H8 02)x
CN
      PMS
PCT
      Polyacrylic
SR
       STN Files:
                        CA, CAPLUS
       CM
             1
       CRN 140-88-5
       CMF C5 H8 02
```

```
CM
     CRN
           80-62-6
          C5 H8 O2
     CMF
                 4 REFERENCES IN FILE CA (1907 TO DATE)
                 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)
    ANSWER 3 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN
     208663-46-1 REGISTRY
RN
     Entered STN: 19 Jul 1998
Acetic acid, 2-propenyl ester, polymer with 1-methylethenyl acetate (9CI)
ED
      (CA INDEX NAME)
OTHER CA INDEX NAMES:
     1-Propen-2-ol, acetate, polymer with 2-propenyl acetate (9CI) (C5 H8 02 . C5 H8 02)\mathbf{x}
MF
     PMS
CI
PCT
     Polyvinyl
SR
     CA
LC
     STN Files:
                    CA, CAPLUS
     CM
     CRN 591-87-7
     CMF C5 H8 O2
AcO-CH_2-CH=-CH_2
           2
     CM
     CRN 108-22-5
     CMF
           C5 H8 O2
     CH<sub>2</sub>
AcO-C-Me
                 1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
                 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L40 ANSWER 4 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN
     121917-49-5 REGISTRY
     Entered STN: 04 Aug 1989
ED
```

L40 ANSWER 4 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN
RN 121917-49-5 REGISTRY
ED Entered STN: 04 Aug 1989
CN 2-Propenoic acid, 2-methyl-, methyl ester, polymer with ethyl
2-propenoate, block (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN 2-Propenoic acid, ethyl ester, polymer with methyl 2-methyl-2-propenoate,
block (9CI)
OTHER NAMES:
CN Ethyl acrylate-methyl methacrylate block copolymer
CN Methyl methacrylate-ethyl acrylate block copolymer

```
(C5 H8 O2 . C5 H8 O2) x
MF
CI
      PMS, COM
PCT
      Polyacrylic
SR
LC
      STN Files:
                      CA, CAPLUS, USPATZ, USPATFULL
      CM
            1
      CRN 140-88-5
      CMF C5 H8 02
      CM
            2
      CRN 80-62-6
           C5 H8 O2
      CMF
  H<sub>2</sub>C
                 29 REFERENCES IN FILE CA (1907 TO DATE)
                 1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
29 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L40 ANSWER 5 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN RN 113547-51-6 REGISTRY ED Entered STN: 26 Mar 1988
      2-Propenoic acid, 2-methyl-, methyl ester, polymer with ethyl 2-propenoate, graft (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      2-Propenoic acid, ethyl ester, polymer with methyl 2-methyl-2-propenoate,
      graft (9CI)
OTHER NAMES:
      Ethyl acrylate-methyl methacrylate graft copolymer (C5 H8 O2 . C5 H8 O2)x
CN
MF
CI
      PMS
      Polyacrylic
PCT
SR
      CA
LC
      STN Files: CA, CAPLUS, USPATFULL
      CM
            1
      CRN 140-88-5
      CMF C5 H8 02
        -CH=CH2
      CM
      CRN 80-62-6
      CMF C5 H8 02
```

```
H<sub>2</sub>C 0
|| ||
Me-C-C-OMe
```

DISPLAY

```
2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
               23 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L40 ANSWER 6 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN
RN
     9010-88-2 REGISTRY
     Entered STN: 16 Nov 1984
ED
     2-Propenoic acid, 2-methyl-, methyl ester, polymer with ethyl 2-propenoate
     (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     2-Propenoic acid, ethyl ester, polymer with methyl 2-methyl-2-propenoate
     Acrylic acid ethyl ester, polymer with methyl methacrylate (8CI)
     Methacrylic acid methyl ester, polymer with ethyl acrylate (8CI)
OTHER NAMES:
CN
     2817D
     Acrilem 674
CN
     Acryloid B 44S
CN
CN
     Acryloid B 72
CN
     Acryloid B 82
CN
     Acryloid K 120N
CN
     Acryloid K 120NL
     Acryloid K 130
Acryloid K 147
CN
CN
CN
     Altuglas 9E
     Altulite 2773
CN
CN
     Cevian A 45000
     CP 41
CN
     CP 41 (acrylate polymer)
Daitosol 5000AD
CN
CN
CN
     Diakon APA 1
     Diakon MG 102
CN
CN
     Dianal BR 64
CN
     Elvacite EX 2612
CN
     Ethyl acrylate-methyl methacrylate copolymer
     Ethyl acrylate-methyl methacrylate polymer
CN
     Eudragit E 30D
CN
     Eudragit NE 30D
CN
     Flexbond 984
CN
     K 120N
CN
     K 130
CN
     K 147
CN
     Kollicoat EMM 30D
CN
     Lucite 30
CN
     Methyl methacrylate-ethyl acrylate copolymer
CN
     Methyl methacrylate-ethyl acrylate polymer
CN
     MG 102
CN
     NeoCryl BT 20
     New Coat PV 400
New Coat PV 412
CN
CN
CN
     Oroglas V 825T
CN
     PA 20
CN
     PA 20 (acrylic polymer)
CN
     Paraloid B 44S
CN
     Paraloid B 72
     Paraloid B 82
CN
CN
     Paraloid K 120N
     Paraloid K 120NL
CN
CN
     Paraloid K 130
CN
     Plexiglas VS 100
     Plexigum MB 319
CN
     Preparation 2817D
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
```

23 REFERENCES IN FILE CA (1907 TO DATE)

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9011-72-7, 9046-85-9, 9048-98-0, 53986-74-6, 54018-07-4, 54183-02-7, 55719-51-2, 60318-38-9, 62534-36-5, 63251-48-9, 97794-17-7, 98036-92-1, 99550-83-1, 51109-48-9, 51311-84-3, 51801-09-3, 52019-30-4, 136303-31-6,
DR
      137087-70-8, 37199-58-9, 72394-19-5, 72626-02-9, 85897-95-6, 30174-68-6, 39301-09-2, 39390-68-6, 107950-48-1, 181591-72-0, 211629-39-9
MF
      (C5 H8 O2 . C5 H8 O2)x
      PMS, COM
CI
PCT
      Polyacrylic
      STN Files: BIOBUSINESS, BIOSIS, CA, CAPLUS, CHEMCATS, CHEMLIST, CIN, CSCHEM, DDFU, DRUGU, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MSDS-OHS, PIRA, PROMT, TOXCENTER, USPAT2, USPATFULL
Other Sources: DSL**, TSCA**
            (**Enter CHEMLIST File for up-to-date regulatory information)
      CM
      CRN 140-88-5
      CMF
            C5 H8 02
        -CH=
             =CH<sub>2</sub>
      CM
      CRN
            80-62-6
      CMF
            C5 H8 O2
**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
                2711 REFERENCES IN FILE CA (1907 TO DATE)
                  55 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
                2714 REFERENCES IN FILE CAPLUS (1907 TO DATE)
=> d fcn 140 6
L40 ANSWER 6 OF 6 REGISTRY COPYRIGHT 2005 ACS on STN
      2-Propenoic acid, 2-methyl-, methyl ester, polymer with ethyl 2-propenoate
       (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
      2-Propenoic acid, ethyl ester, polymer with methyl 2-methyl-2-propenoate
CN
      Acrylic acid ethyl ester, polymer with methyl methacrylate (8CI)
      Methacrylic acid methyl ester, polymer with ethyl acrylate (8CI)
OTHER NAMES:
CN
      2817D
      Acrilem 674
CN
CN
      Acryloid B 44S
      Acryloid B 72
CN
      Acryloid B 82
CN
      Acryloid K 120N
      Acryloid K 120NL
      Acryloid K 130
CN
      Acryloid K 147
CN
      Altuglas 9E
      Altulite 2773
CN
CN
      Cevian A 45000
CN
      CP 41
```

CP 41 (acrylate polymer)

```
Daitosol 5000AD
     Diakon APA 1
CN
     Diakon MG 102
     Dianal BR 64
     Elvacite EX 2612
CN
     Ethyl acrylate-methyl methacrylate copolymer
CN
      Ethyl acrylate-methyl methacrylate polymer
     Eudragit É 30D
CN
CN
CN
CN
CN
CN
CN
      Eudragit NE 30D
     Flexbond 984
     K 120N
     K 130
     K 147
      Kollicoat EMM 30D
      Lucite 30
CN
CN
CN
      Methyl methacrylate-ethyl acrylate copolymer
     Methyl methacrylate-ethyl acrylate polymer
      MG 102
CN
CN
     NeoCryl BT 20
     New Coat PV 400
CN
CN
     New Coat PV 412
Oroglas V 825T
CN
     PA 20
CN
     PA 20 (acrylic polymer)
     Paraloid B 44S
Paraloid B 72
CN
CN
CN
     Paraloid B 82
     Paraloid K 120N
Paraloid K 120NL
CN
CN
CN
     Paraloid K 130
     Plexiglas VS 100
CN
CN
CN
CN
CN
CN
     Plexigum MB 319
     Preparation 2817D
     Primal AC 22
Primal AC 33
     Primal B 52
     Rhoplex AC 22
Rhoplex AC 33
CN
CN
      Rhoplex AC 35
CN
CN
     Rhoplex B 52
CN
      Romacril ER
CN
      Rowney Cryla
      Sokrat 602
CN
CN
CN
      Sumipex EX
      Sumipex LG 35
CN
      Ucecryl H
CN
      VS 100
      Yodo Sol GH 28
```

# => d his

(FILE 'HOME' ENTERED AT 07:56:46 ON 19 APR 2005)

# FILE 'REGISTRY' ENTERED AT 07:57:20 ON 19 APR 2005 ACT QAZ194CEP/A

]	L1		QUE	ABB=ON PLU=	ON (PMS	OR MAN	OR IDS)/CI OR UNSPECIFIE
]	L2	(	79) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	C16H16N2O6S2 AND SC4/ES AND N
]	L3	(	43) SEA	FILE=REGISTRY	ABB=ON	PLU=ON	L2 AND NR=3
]	[4	(	35) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	L3 NOT (MXS/CI OR MIXT OR MET
]	L5	(	29) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	(1051-68-9/BI OR 117091-43-7/
]	L6	(	1) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	L5 AND 3-METHYL
]	լ7	(	28) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	L5 NOT L6
]	L8	(	32) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	C18H19N3O5S AND NC3-NCSC3/ES
]	L9	(	26) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	L8 NOT L1
]	L10	Ì	5) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	L9 AND CEFPROZ?
	LII	Ì		FILE=REGISTRY		PLU=0N	C25H28N6O7S3 AND NR=4 AND NC3
	L12	Ì	11) SEA	FILE=REGISTRY	ABB=ON	PLU=0N	L11 NOT NC5/ES
	L13	(		FILE=REGISTRY		PLU=0N	C16H17N3O5S AND NR=3 AND NC3-

Qazi 10/642194 Page 82

```
2) SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
                                                        C21H27N509S2 AND NR=3 AND NC3
L14 (
               2) SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
                                                        C20H22N4O10S AND NR=3 AND NC3
L15
               4) SEA FILE=REGISTRY ABB=ON
                                                        C15H14CLN3O4S AND CEFAC? AND
                                               PLU=0N
L16
L17
               5) SEA FILE=REGISTRY ABB=ON
                                               PLU=0N
                                                        C18H18N6O5S2 AND CEFAMAND? AN
                                               PLU=0N
                                                        L17 NOT COMPD
               4) SEA FILE=REGISTRY ABB=ON
L18
L19
              17) SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
                                                        C16H17N3O7S2 AND NR=3 AND NC3
              17) SEA FILE=REGISTRY ABB=ON
                                                        C17H17N3O6S2 AND NR=3 AND NC3
                                               PLU=0N
L20
                                                        (21593-22-6/BI OR 21593-23-7/
L21
              15) SEA FILE=REGISTRY ABB=ON
                                               PLU=0N
L22
              44) SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
                                                        C13H13N5O5S2 AND NCSC2/ES AND
L23
              35) SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
                                                        L22 NOT L1
                                                        L23 NOT (MXS/CI OR MIXT)
              28) SEA FILE=REGISTRY ABB=ON
L24
                                               PLU=0N
                                               PLU=ON
                                                        (102044-69-9/BI OR 109323-67-
L25
              18) SEA FILE=REGISTRY ABB=ON
               8) SEA FILE=REGISTRY ABB=ON
6) SEA FILE=REGISTRY ABB=ON
                                                        C18H18N608S3 AND N4C/ES AND C
L26
                                               PLU=ON
                                               PLU=ON
                                                        L26 NOT MIXT
L27
               5) SEA FILE=REGISTRY ABB=ON
                                               PLU=0N
                                                        L27 NOT COMPD
L28
                                               PLU=0N
L29
             118 SEA FILE=REGISTRY ABB=ON
                                                        L7 OR L10 OR L12 OR L13 OR L1
                  ACT QAZ194GAL1/A
                                                        "XANTHAN GUM"/CN
               1) SEA FILE=REGISTRY ABB=ON PLU=ON
L30 (
                                                        "GUAR GUM"/CN
               1) SEA FILE=REGISTRY ABB=ON
                                               PLU=0N
L31 (
               2 SEA FILE=REGISTRY ABB=ON
                                               PLU=0N
                                                        L30 OR L31
L32
                  ACT QAZ194GAL2/A
             100) SEA FILE=REGISTRY ABB=ON PLU=ON
                                                        XANTHAN (1A) GUM?
L33 (
                                                        L33 NOT (COMPD OR COMPOUND)
              82) SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
L34
L35
              53) SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
                                                        L34 NOT PMS/CI
             290) SEA FILE=REGISTRY ABB=ON
                                               PLU=0N
                                                        GUAR (W) GUM
L36
                                               PLU=0N
             288) SEA FILE=REGISTRY ABB=ON
                                                        L36 NOT (COMPD OR COMPOUND)
L37
L38
             338 SEA FILE=REGISTRY ABB=ON
                                               PLU=ON
                                                        L35 OR L37
                  ACT QAZ194P0L1/A
              32) SEA FILE=REGISTRY ABB=ON PLU=ON "(C5H8O2. C5H8O2) X"/MF
L39 (
               6 SEA FILE=REGISTRY ABB=ON PLU=ON
                                                        (113547-51-6/BI OR 121917-49-
L40
     FILE 'HCAPLUS' ENTERED AT 08:05:34 ON 19 APR 2005
L41
           11259 L29 OR CEFADROXIL OR CEFTIZOXIME OR CEFPODOXIME (1A) PROXET? OR
             521 BMY28167 OR BMY (1A) (28167 OR 28 (1A) 167) OR CEFZIL# OR PRO!E
L42
            1813 CEFODIE OR MONOCIDUR# OR MOPNOCID# OR PRATICEF# OR CEFIZOX# OR
L43
            286 CEFTISOMIN# OR EPOCELIN# OR DURICEF# OR CCI15641 OR CCI (1A) (1 122 LILLY99368 OR LILLY (1A) (99368 OR 99 (1A) 368) OR PANORAL# OR 1207 HYDROXYCEPHALIN# OR CEPHITOXIN# OR CEFADOLE OR CEFAMANDOL OR CE
L44
L45
L46
             525 CEFADYL# OR CEFALOJECT# OR CEFAPRIN# OR CEFATRAXYL OR CEPHATREX
L47
            2500 CEPHALOSPORAN? (1A) ACID OR CEPHALOSPORANATE OR ?ACETAMIDO (1A) 92 MICROTIN# OR SEFFIN# OR ?ACETAMIDO/BI (1A) CEPH OR SYNCLOTIN# O
L48
L49
                  E CEPHALEXIN/CT
                  E E3+ALL
L50
            3267 CEPHALEXIN/CT
                  E CEFPROZIL/CT
                  E CEFDITOREN/CT
                  E CEFADROXIL/CT
                  E E3+ALL
             881 CEFADROXIL/CT
L51
                  E CEFPODOXIME/CT
                  E CEFUROXIME/CT
                  E E3+ALL
            2607 CEFUROXIME/CT
L52
                  E CEFACLOR/CT
                  E E3+ALL
            1483 CEFACLOR/CT
L53
                  E CEFAMANDOLE/CT
                  E E3+ALL
            1642 CEFAMANDOLE/CT
L54
                  E CEFITOXIN/CT
                  E CEFOXITIN/CT
                  E E3+ALL
L55
            3075 CEFOXITIN/CT
```

```
E CEPALOTHIN/CT
                     E CEPHAPIRIN/CT
                     E E3+ALL
L56
               561 CEPHAPIRIN/CT
                     E CEFTIZOXIME/CT
                     E E3+ALL
              1070 CEFTIZOXIME/CT
L57
                    E CEFONICID/CT
       FILE 'REGISTRY' ENTERED AT 09:17:12 ON 19 APR 2005
                     E GALACTOMANNAN/CN
                 75 GALACTOMANNAN OR GALACTO (1A) MANNAN?
L58
       FILE 'HCAPLUS' ENTERED AT 09:18:24 ON 19 APR 2005
              3355 L58 OR GALACTO (1A) MANNAN? OR DAIKOL? OR FAIBARON? OR GALACTOM
L59
                    E GALCTOMANNAN/CT
                    E GALACTOMANNAN/CT
                    E E3+ALL
L60
              2078 GALACTOMANNAN+OLD/CT
             49966 L32 OR L38 OR XANTHAN (1A) GUM OR ACTIGUM? OR ADM40 OR ADM (1A)
L61
               893 IDVIS OR INAGEL? OR JUNGBUNZLAUER? OR K5C151 OR K9C57 OR K (1A)
L62
L63
             38695 GUAR (1A) GUM OR EMCOGUM? OR DEALCA? OR EMULGUM? OR FINE (1A) G
                    E XANTHAN/CT
                     E E4+ALL
L64
              8814 XANTHAN GUM/CT
                     E GUAR GUM/CT
                    E E3+ALL
L65
              7040 GUAR GUM+NT/CT
                    E LOCUAT/CT
                    E BEAN/CT
                    E BEAN GUM/CT
                    E E4+ALL
                    E SWELLING AGENTS/CT
                    E E3+ALL
L66
               105 SWELLING AGENTS/CT (L) POLYMER?
              5193 (POLYMER?/CW OR POLYMERS+OLD, NT1/CT) (L) SWELL?
L67
             7289 L40 OR PROPENOIC (IA) ACID (IA) 2 (IA) METHYL (5A) ETHYL (IA) P 10274 ALTUGLAS? OR ALTIVITE? OR CEVIAN? OR CP41? OR CP (IA) 41 OR DAI 2505 NEOCRYL? OR NEW (IA) COAT (IA) (PV400 OR PV412 OR PV (IA) (400
L68
L69
L70
              6989 ROWNEY (1A) CRYLA OR SOKRAT? OR SUMIPEX? OR UCECRYL? OR VS100 O E KSHIRSAGAR R/AU
L71
L72
                 10 E3, E6-7
                    E BOLDHANE S/AU
L73
                  3 E4
                    E JINDAL K/AU
L74
                 34 E3-4, E10
                64 (ORCHID AND CHEM? AND PHARM?)/CS, PA
L75
L76
                    QUE PY<=2003 OR AY<=2003 OR PRY<=2003 OR PD<20030818 OR AD<2003
L77
                58 L41-57 AND L58-65
L78
                 5 L77 AND L72-75
L79
                53 L77 NOT L78
L80
                 6 L79 AND L68-71
L81
                 6 L80 AND L76
=> b hcap
FILE 'HCAPLUS' ENTERED AT 09:47:58 ON 19 APR 2005
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DATE

Qazi 10/642194 FILE COVERS 1907 - 19 Apr 2005 VOL 142 ISS 17 FILE LAST UPDATED: 18 Apr 2005 (20050418/ED) New CAS Information Use Policies, enter HELP USAGETERMS for details. This file contains CAS Registry Numbers for easy and accurate substance identification. => d all 178 tot L78 ANSWER 1 OF 5 HCAPLUS COPYRIGHT 2005 ACS on STN AN 2004:412577 HCAPLUS DN 140:395542 Entered STN: 21 May 2004 ED Sustained release pharmaceutical composition of a cephalosporin antibiotic ΤI Kshirsagar, Rajesh Suresh; Boldhane, Sanjay Parbhatrao IN ; Jindal, Kour Chand PA Orchid Chemicals & Pharmaceuticals Limited, India U.S. Pat. Appl. Publ., 21 pp., Cont.-in-part of U.S. Ser. No. 222, 930. S<sub>0</sub> CODEN: USXXCO DT Patent English LA ICM A61K031-545 IC ICS A61K009-20; A61K031-736 424465000; 514200000; 514054000 NCL 63-6 (Pharmaceuticals) FAN. CNT 3 PATENT NO. KIND DATE APPLICATION NO. PΙ US 2004096496 A1 20040520 US 2003-642194 US 2004033262 20040219 US 2002-222930 A1 WO 2002-IB3320 20040226 WO 2004016250 A1 W: AE, AG, AL, AM, AT, AU, AZ, DZ, EC, EE, ES, CU, CZ, DE, DK, DM, CO, CR, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, . KP, LV. LU, MA, MD, MG, MK, MN, MW, MX, LS, LT, PL, PT, R0, RU, SD, SE, SG, SI, SK, SL, TJ, VC, VN, YU, ZA, ZM, ZW UA, UG, US, UZ, UG, TZ,

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AB Sustained release pharmaceutical compns. comprising at least a cephalosporin antibiotic, a mixture of polymers comprising of galactomannans and neutral swellable polymers, and other pharmaceutically acceptable excipients are described. The composition comprises about 30% to about 90% by weight of a cephalosporin antibiotic; about 1% to about 30% by weight of said mixture of polymers comprising from about 0.1% to about 15% by weight of galactomannans, and about 0.1% to about 15% of neutral swellable polymer by weight of sustained release composition

cephalosproin antibiotic sustained release tablet ST

IT Drug delivery systems

(carriers; sustained-release pharmaceutical composition of a cephalosporin antibiotic)

IT Lubricants

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(pharmaceutical; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
ΙT
     Swelling agents
        (polymers; sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
IT
    Antibiotics
     Binders
     Drying
     Mixing
    Plasticizers
     Sieving
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
    Gelatins, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     Polyesters, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
IT
    Drug delivery systems
        (sustained-release; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
     Polymers, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (swellable; sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
IT
    Drug delivery systems
        (tablets, coated; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
IT
     Drug delivery systems
        (tablets; sustained-release pharmaceutical composition of a cephalosporin
        antibiotic)
     Fats and Glyceridic oils, biological studies
IT
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (vegetable, hydrogenated; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
     7631-86-9, Silicon dioxide, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
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        (colloidal; sustained-release pharmaceutical composition of a cephalosporin
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     9004-34-6, Cellulose, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (microcryst.; sustained-release pharmaceutical composition of a
        cephalosporin antibiotic)
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     9005-25-8, Starch, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
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        (pregelatinized; sustained-release pharmaceutical composition of a
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    9003-39-8, Polyvinylpyrrolidone 9
9004-64-2, Hydroxypropyl cellulose
                                        9004-62-0, Hydroxyethylcellulose
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                                          9004-65-3, Hydroxypropyl
     methylcellulose
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     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     57-11-4, Stearic acid, biological studies 57-50-1, Sucrose, biological
                                              557-05-1, Zinc stearate
     studies
              557-04-0, Magnesium stearate
     1592-23-0, Calcium stearate 9000-30-0, Guar gum 11138-66-2, Xanthan gum
     14807-96-6, Talc, biological studies
     RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     153-61-7, Cephalothin 11078-30-1, Galactomannan
     11111-12-9D, Cephalosporins, derivs. 15686-71-2, Cephalexin
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21593-23-7, Cephapirin 34444-01-4,
     Cefamandole 35607-66-0, Cefoxitin 50370-12-2, Cefadroxil 53994-73-3, Cefaclor
     61270-58-4, Cefonicid 64544-07-6, Cefuroxime
     axetil 68401-81-0, Ceftizoxime 87239-81-4,
     Cefpodoxime proxetil 92665-29-7,
     Cefprozil 117467-28-4, Cefditoren
     pivoxil
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); THU (Therapeutic use); BIOL (Biological study); PROC (Process);
     USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     9004-32-4
                  9010-88-2, Ethyl acrylate-methyl methacrylate copolymer
     9050-04-8, Calcium carboxymethyl cellulose 25086-89-9, Plasdone S-630 RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
     50-70-4, Sorbitol, biological studies 50-99-7, Glucose, biological studies 63-42-3, Lactose 69-65-8, Mannitol 69-79-4, Maltose
     9004-53-9, Dextrin
                                                      74811-65-7, Croscarmellose
                            66828-18-0, Dextrate
     sodium
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained-release pharmaceutical composition of a cephalosporin antibiotic)
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     2004:162583 HCAPLUS
     140:205146
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     Entered STN: 29 Feb 2004
     Sustained release pharmaceutical compositions of a cephalosporin
ΤI
     Kshirsagar, Rajesh Suresh; Boldhane, Sanjay Parbhatrao
IN
      ; Jindal, Kour Chand
     Orchid Health Care, A Division of Orchid
PA
     Chemicals & Pharmaceuticals Ltd., India
S0
     PCT Int. Appl., 55 pp.
     CODEN: PIXXD2
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     English
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     ICS A61K031-545; A61K031-546
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                           A61K009/20H6B; A61K009/20H6F; A61K031/545; A61K031/546;
                           A61K031/736
     Sustained release pharmaceutical compns. comprising at least a
     cephalosporin, a mixture of polymers comprising of galactomannans
     and neutral swellable polymers, and other pharmaceutically acceptable
     excipients are described. The composition comprises 30-90% of a cephalosporin
     antibiotic, 1-30% mixture of polymers comprising 0.1-15%
     galactomannans, and 0.1-15% by weight of a neutral swellable polymer.
     Thus, tablets contained cephalexin monohydrate 75.73, lactose 6.27,
     xanthan gum 7.0, Eudragit NE 30D 7.0, HPMC E5 31.5, Mg
     stearate 1.0 and water qs to 100%.
     sustained release cephalosporin polymer
     Drug delivery systems
         (granules, sustained release; sustained release pharmaceutical compns.
         of cephalosporins)
IT
     Binders
     Dissolution
     Drug bioavailability
     Human
     Lubricants
     Plasticizers
         (sustained release pharmaceutical compns. of cephalosporins)
     Carbohydrates, biological studies
     Gelatins, biological studies
Polymers, biological studies
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained release pharmaceutical compns. of cephalosporins)
     Drug delivery systems
ΙT
         (sustained-release; sustained release pharmaceutical compns. of
         cephalosporins)
IT
     Drug delivery systems
         (tablets, sustained-release; sustained release pharmaceutical compns.
         of cephalosporins)
     Fats and Glyceridic oils, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (vegetable, hydrogenated; sustained release pharmaceutical compns. of
         cephalosporins)
     9004-34-6, Cellulose, biological studies
ΙT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (microcryst.; sustained release pharmaceutical compns. of
         cephalosporins)
     15686-71-2, Cephalexin 50370-12-2, Cefadroxil
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     RL: PKT (Pharmacokinetics); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sustained release pharmaceutical compns. of cephalosporins)
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         (sustained release pharmaceutical compns. of cephalosporins)
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(1) Lupin Lab Ltd; WO 0241876 A 2002 HCAPLUS
   Shankar, M; WO 02067943 A 2002 HCAPLUS
(3) Squibb Bristol Myers Co; WO 9846213 A 1998 HCAPLUS
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     Sustained-release pharmaceutical composition of a cephalosporin antibiotic
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     Kshirsagar, Rajesh Suresh; Boldhane, Sanjay Parbhatrao
      ; Jindal, Kour Chand
     Orchid Health Care, India
U.S. Pat. Appl. Publ., 10 pp.
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BJ, CF,
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                      GB,
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                                  20040520
     US 2004096496
                                               US 2003-642194
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                            A1
PRAI IN 2002-MA601
                                  20020816
                            A
        2002-222930
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                            Α
                                  20020819
        2002-IB3320
     WO
                            Α
CLASS
                  CLASS PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
 US 2004033262
                  ICM
                          A61K031-545
                          A61K031-736; A61K009-20; A61K009-22
                  ICS
                          424468000; 514054000; 514200000
                  NCL
                         A61K009/20H6B; A61K009/20H6F; A61K031/545; A61K031/546;
 US 2004033262
                  ECLA
                         A61K031/736
                         A61K031/545; A61K031/546
                  ECLA
 US 2004096496
     A sustained release pharmaceutical composition comprising at least a
     cephalosporin antibiotic and a mixture of polymers and other
     pharmaceutically acceptable excipients is described. Polymers are
     selected from mixture of galactomannans and neutral swellable
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polymers, which releases the active ingredient in a predetd. manner. For
     example, a sustained-release tablet composition contained cephalexin 795.32 mg,
     lactose 107.68 mg, xanthan gum 31.5 mg, Eudragit NE 30D 52.5 mg, HPMC E5 52.5 mg, and magnesium stearate 10.5 mg. Cephalexin release from the tablets was 25.21%, 50.84%, 73.18%, and 84.17% after 1h,
                                                                         Cephalexin
     6 h, 10 h and 14 h, resp.
     cephalosporin polymer mixt sustained release tablet
     Dissolution
     Drug bioavailability
     Granulation
        (sustained-release composition of cephalosporin antibiotic containing mixture of
        polymers)
IT
     Polymers, biological studies
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (sustained-release composition of cephalosporin antibiotic containing mixture of
        polymers)
IT
     Drug delivery systems
        (sustained-release; sustained-release composition of cephalosporin
        antibiotic containing mixture of polymers)
IT
     Drug delivery systems
        (tablets, sustained-release; sustained-release composition of cephalosporin
        antibiotic containing mixture of polymers)
     9000-30-0, Guar gum 9000-40-2, Locust bean
     gum 9010-88-2, Eudragit NE 30D 11078-30-1, Galactomannan 11138-66-2, Xanthan gum
     RL: POF (Polymer in formulation); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (sustained-release composition of cephalosporin antibiotic containing mixture of
        polymers)
     153-61-7, Cephalothin 11111-12-9, Cephalosporin
     15686-71-2, Cephalexin 21593-23-7, Cefaprin
     34444-01-4, Cefamandole 35607-66-0, Cefoxitin 50370-12-2, Cefadroxil 53994-73-3,
     Cefaclor 61270-58-4, Cefonicid
     64544-07-6, Cefuroxime axetil 68401-81-0,
     Ceftizoxime 87239-81-4, Cefpodoxime
     proxetil 92665-29-7, Cefprozil
     117467-28-4, Cefditoren pivoxil
     RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
        (sustained-release composition of cephalosporin antibiotic containing mixture of
        polymers)
     ANSWER 4 OF 5 HCAPLUS COPYRIGHT 2005 ACS on STN
L78
AN
     2003:417601 HCAPLUS
DN
     139:12261
ED
     Entered STN: 01 Jun 2003
     A pharmaceutical composition for controlled release of a β-lactam
ΤI
     antibiotic
     Sen, Himadri; Kshirsagar, Rajesh Suresh; Bhamare, Shailesh
IN
     Suresh
     Lupin Limited, India
     PCT Int. Appl., 36 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
IC
     ICM A61K009-22
     ICS A61K031-545; A61K031-24
     63-6 (Pharmaceuticals)
     Section cross-reference(s): 1
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     PATENT NO.
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              GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
              LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO,
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              RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US,
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Qazi 10/642194

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LU, MC, NL, PT,
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                                             BR 2001-17171
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     BR 2001017171
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     US 2005031685
                                             US 2004-495961
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PRAI WO 2001-IN204
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CLASS
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 WO 2003043607
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                        A61K031-545; A61K031-24
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                         A61K009/20H2; A61K009/20H4B; A61K009/20H6F;
 US 2005031685
                 ECLA
                        A61K009/20H6F2; A61K009/20K4B; A61K031/24;
                         A61K031/24+M; A61K031/545; A61K031/545+M; A61K033/06+M
     An improved stable pharmaceutical composition for controlled release of an
AB
     active ingredient comprises a β-lactam antibiotic such as cephalexin,
     cefaclor or their hydrates, salts or esters as active ingredients,
     a calcium salt and a mixture of hydrophilic polymers selected from the group
     consisting of sodium alginate and xanthan gum and with
     or without hydroxypropyl Me cellulose, the composition optionally containing
     probenecid. The composition may also contain a water soluble and/or water
     dispersible diluent, wherein the quantities of the hydrophilic polymers
     and water soluble and/or water dispersible diluents are such that the active
     ingredient is released at a rate suitable for once or twice daily
     administration of the pharmaceutical composition Thus, tablets contained
     cephalexin 73.87, sodium alginate 6.00, xanthan gum
     5.00, Methocel K15M 5.00, calcium sulfate 0.72, lactose monohydrate 7.40,
     and Mg stearate 2.00% by weight
     pharmaceutical controlled release beta lactam antibiotic
     Drug delivery systems
IT
        (controlled-release; pharmaceutical composition for controlled release of
        β-lactam antibiotics)
IT
     Dissolution
     Drug bioavailability
     Human
     Lubricants
        (pharmaceutical composition for controlled release of β-lactam
        antibiotics)
    Polymers, biological studies
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
IT
        (pharmaceutical composition for controlled release of β-lactam
        antibiotics)
     Drug delivery systems
IT
        (solids, controlled release; pharmaceutical composition for controlled
        release of β-lactam antibiotics)
ΙT
     Drug delivery systems
        (tablets, controlled-release; pharmaceutical composition for controlled
        release of β-lactam antibiotics)
IT
     Antibiotics
        (β-lactam; pharmaceutical composition for controlled release of
        β-lactam antibiotics)
IT
     9004-34-6, Cellulose, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (microcryst.; pharmaceutical composition for controlled release of
        β-lactam antibiotics)
     15686-71-2, Cephalexin
IT
     RL: PKT (Pharmacokinetics); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (pharmaceutical composition for controlled release of \beta-lactam
        antibiotics)
                                     63-42-3, Lactose
                                                          299-28-5, Calcium
                           62-33-9
     57-66-9, Probenecid
                 557-04-0
                            814-80-2, Calcium lactate
                                                          5743-27-1, Calcium
     gluconate
                 7693-13-2, Calcium citrate 7778-18-9, Calcium sulfate
     ascorbate
                                                  9005-38-3, Sodium alginate
     9004-65-3, Hydroxypropyl methyl cellulose
     9063-38-1, Starch, carboxymethyl ether, sodium salt 10043-52-4, Calcium
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chloride, biological studies 11138-66-2, Xanthan
     gum 53994-73-3, Cefaclor 64044-51-5, Lactose
     monohydrate
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (pharmaceutical composition for controlled release of β-lactam
        antibiotics)
              THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE. CNT 4
RE
   Boots Co Plc; EP 0234670 A 1987 HCAPLUS
(1)
   Katz; WO 9822091 A 1998 HCAPLUS
    Staniforth, J; WO 0015198 A 2000 HCAPLUS
(4) Toyama Chem Co Ltd; JP 52105220 A 1977 HCAPLUS
    ANSWER 5 OF 5 HCAPLUS COPYRIGHT 2005 ACS on STN
L78
     2002:408511 HCAPLUS
     136:406868
DN
ED
     Entered STN: 31 May 2002
     Pharmaceutical composition for controlled release of a β-lactam
     Sen. Himadri; Kshirsagar, Rajesh S.; Kandi, Chandrashekhar S.;
IN
     Bhamare, Shailesh
PA
     Lupin Laboratories Limited, India
SO
     PCT Int. Appl., 38 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
     ICM A61K009-22
IC
     ICS A61K009-24; A61K031-545
CC
     63-6 (Pharmaceuticals)
FAN. CNT 1
                                             APPLICATION NO.
                                                                     DATE
     PATENT NO.
                          KIND
                                 DATE
                                             WO 2000-IN112
                                                                     20001122
     WO 2002041876
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     US 2002103181
                                 20020801
                                             US 2000-726636
                                                                     20001130
                           A1
PRAI WO 2000-IN112
                                 20001122
CLASS
                        PATENT FAMILY CLASSIFICATION CODES
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                 ICM
                         A61K009-22
 WO 2002041876
                         A61K009-24; A61K031-545
                  ICS
                         A61K009/20H6F; A61K009/20K4B; A61K031/545
                 ECLA
 US 2002103181
     A tablet for controlled release of a β-lactam antibiotic, such as
     cephalexin, cefaclor or their pharmaceutically acceptable
     hydrates, salts or esters, comprises a mixture of hydrophilic polymers
     selected from the group consisting of at least one sodium alginate and at
     least one xanthan gum as controlled release matrix;
     and optionally probenecid as an antibiotic adjuvant as either immediate
     release or controlled release part. The composition may also contain one or
     more of a water soluble and/or water dispersible diluent, wherein the
     quantities of the hydrophilic polymer matrix still provides the desired
     once a day profile. The resulting modified release matrix formulation not
     containing probenecid may be administered once or twice daily, while the
     formulation containing probenecid may be administered once daily. For
     example, cephalexin controlled-release tablets were prepared from two parts:
     (1) a controlled-release part containing (by weight) cephalexin 72.58%, sodium
     alginate 14.00%, xanthan gum 7.00%, mannitol 5.45%,
     and magnesium stearate 0.96%, and an immediate-release part containing
     probenecid 79.36%, microcryst. cellulose 15.15%, sodium starch glycolate
     5.00%, and magnesium stearate 0.47%. Cephalexin release in simulated
     intestinal fluid without pancreatin was 28.0%, 61.80%, 92.70%, and 101.60%
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after 1, 4, 10, and 12 h, while probenecid release was 98.80%, 102.20%, and 103.30% after 10, 20, and 30 min, resp.
     beta lactam antibiotic controlled release tablet dissoln
    Polymers, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (hydrophilic; preparation and dissoln. of β-lactam antibiotic
        controlled-release tablets)
     Compaction
     Compression
     Dissolution
         (preparation and dissolm. of β-lactam antibiotic controlled-release
        tablets)
IT
    Biopolymers
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (preparation and dissoln. of β-lactam antibiotic controlled-release
        tablets)
    Drug delivery systems
IT
         (tablets, controlled-release; preparation and dissoln. of β-lactam
        antibiotic controlled-release tablets)
IT
    Lactams
     RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
        (\beta-, antibiotics; preparation and dissoln. of \beta-lactam antibiotic
        controlled-release tablets)
     Antibiotics
IT
         (\beta-lactam; preparation and dissoln. of \beta-lactam antibiotic
        controlled-release tablets)
     7631-86-9, Colloidal silicon dioxide, biological studies
IT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (colloidal, disintegrant; preparation and dissoln. of β-lactam
        antibiotic controlled-release tablets)
     63-42-3, Lactose RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
ΙT
         (diluent; preparation and dissoln. of β-lactam antibiotic
        controlled-release tablets)
     9004-34-6, Cellulose, biological studies
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
IT
         (microcryst., diluent; preparation and dissoln. of β-lactam antibiotic
        controlled-release tablets)
     57-66-9, Probenecid RL: MOA (Modifier or additive use); THU (Therapeutic use); BIOL
IT
     (Biological study); USES (Uses)
         (preparation and dissolm. of \beta-lactam antibiotic controlled-release
        tablets)
     15686-71-2, Cephalexin 53994-73-3, Cefaclor
     RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES
     (Uses)
         (preparation and dissoln. of β-lactam antibiotic controlled-release
        tablets)
                           557-04-0, Magnesium stearate 9005-38-3, Sodium alginate 9
                                                              9005-25-8, Starch,
     69-65-8, D-Mannitol
                                                          9063-38-1, Sodium starch
     biological studies
     glycolate 11138-66-2, Xanthan gum
     14807-96-6, Talc, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (preparation and dissoln. of β-lactam antibiotic controlled-release
        tablets)
               THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE. CNT 3
RE
(1) The Boots Company Plc; EP 0234670 A 1987 HCAPLUS
    Toyama Chem Co Ltd; JP 52105220 A 1977 HCAPLUS
(3) Yissum Research Development Company; WO 9822091 A 1998 HCAPLUS
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L81 ANSWER 1 OF 6 HCAPLUS COPYRIGHT 2005 ACS on STN

AN 2003:818128 HCAPLUS

DN 139:311996

ED Entered STN: 17 Oct 2003

TI Targeted controlled delivery compositions activated by changes in pH or

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salt concentration
     Shefer, Adi; Shefer, Shmuel David
IN
PA
     U.S. Pat. Appl. Publ., 20 pp.
SO
     CODEN: USXXCO
DT
     Patent
LA
     English
IC
      ICM A61K007-50
     510444000; 510130000; 510504000
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      62-5 (Essential Oils and Cosmetics)
      Section cross-reference(s): 46, 63
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      WO 2003087287
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                                    20031023
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PRAI US 2002-119567
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      WO 2003-US9607
CLASS
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 PATENT NO.
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                           A61K007-50
 US 2003195133
                           510444000; 510130000; 510504000
                   NCL
                           A61K008/04H; A61Q013/00; C11D003/00B9; C11D003/50B2;
 US 2003195133
                   ECLA
                           C11D017/00D
     The present invention relates to a novel controlled release carrier system
AR
      for pH or salt triggered release and targeted delivery of fragrances and
      other active ingredients onto fabric, hair, skin, and other biol. surfaces
      and which provides prolonged release of fragrances and other active
     ingredients over an extended period of time, or yields a high impact fragrance "burst" upon treating the target surface with heat (blow drying
      the hair, ironing the fabric). The controlled delivery system of the
      present invention is substantially a free-flowing powder formed of solid
      hydrophobic nanospheres comprising the fragrance and other active
      ingredients that are encapsulated in a pH- or salt-sensitive microspheres.
      Also described are processes for preparing such compns. and processes for
      using same. Furthermore, certain components of the aforementioned compns.
      in combination with one another are novel, and other components have novel
      uses in increasing fragrance substantivity, particularly in fabric, hair,
      and skin care prepns. The invention further pertains to consumer and
      diversified products comprising the controlled release system of the
     present invention. For example, polyethylene was melted in an oven, then mixed with water, Eudragit EPO, Hi-Cap 100, and
     polyethyleneimine. To the mixture, Incroquat Behenyl HE and a fragrance solution were added to form a suspension. The suspension was spray-dried to produce a free flowing, dry powder, consisting of 30 % fragrance
      encapsulated in solid hydrophobic nanospheres. Shampoos containing the above
      fragrances deposited higher level of fragrances on the hair, as compared
      to the control samples comprising the neat fragrances.
      controlled release pH salt sensitive microsphere; fragrance encapsulation
ST
      hydrophobic polymer shampoo
     Fats and Glyceridic oils, biological studies
RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
      (Biological study); USES (Uses)
          (Japan wax; targeted controlled delivery microspheres activated by
         changes in pH or salt concentration)
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Phenols, biological studies RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
IT
     (Biological study); USES (Uses)
        (alkyl, ethoxylated; targeted controlled delivery microspheres
        activated by changes in pH or salt concentration)
     Phenols, biological studies RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
IT
     (Biological study); USES (Uses)
         (alkyl; targeted controlled delivery microspheres activated by changes
        in pH or salt concentration)
IT
     Surfactants
         (cationic; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
     Polyoxyalkylenes, biological studies
IT
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (di-Me, Me hydrogen polysiloxane-; targeted controlled delivery
        microspheres activated by changes in pH or salt concentration)
     Polysiloxanes, biological studies
ΙT
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (di-Me, Me hydrogen, polyoxyalkylene-; targeted controlled delivery
        microspheres activated by changes in pH or salt concentration)
     Quaternary ammonium compounds, biological studies
IT
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (dicoco alkyldimethyl, Me sulfates; targeted controlled delivery
        microspheres activated by changes in pH or salt concentration)
     Quaternary ammonium compounds, biological studies
RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
IT
     (Biological study); USES (Uses)
         (dimethylditallow alkyl, Me sulfates; targeted controlled delivery
        microspheres activated by changes in pH or salt concentration)
     Quaternary ammonium compounds, biological studies
IT
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (dimethylditallow alkyl, chlorides; targeted controlled delivery
        microspheres activated by changes in pH or salt concentration)
     Fatty acids, biological studies
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (esters; targeted controlled delivery microspheres activated by changes
        in pH or salt concentration)
IT
     Monoglycerides
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (ethoxylated; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
     Alcohols, biological studies
ΙT
     Amines, biological studies
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (fatty, ethoxylated; targeted controlled delivery microspheres
        activated by changes in pH or salt concentration)
     Alcohols, biological studies RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
IT
     (Biological study); USES (Uses)
         (fatty; targeted controlled delivery microspheres activated by changes
         in pH or salt concentration)
IT
     Gelatins, biological studies
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (formalized; targeted controlled delivery microspheres activated by
         changes in pH or salt concentration)
IT
     Surfactants
         (ionic; targeted controlled delivery microspheres activated by changes
         in pH or salt concentration)
     Drug delivery systems
IT
         (microspheres; targeted controlled delivery microspheres activated by
         changes in pH or salt concentration)
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IT
     Cosmetics
        (moisturizers; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
IT
     Surfactants
        (nonionic; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
     Polyamines
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (polyalkylene-; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
     Alcohols, biological studies
IT
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (polyhydric; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
IT
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (sandarac; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
IT
     Anesthetics
     Anti-inflammatory agents
     Antibacterial agents
     Antibiotics
     Antioxidants
     Fabric softeners
     Gums and Mucilages
     Hair preparations
     Hydrocolloids
     Odor and Odorous substances
     Perfumes
     Sunscreens
        (targeted controlled delivery microspheres activated by changes in pH
        or salt concentration)
IT
     Candelilla wax
     Glutens
     Keratins
     Paraffin oils
     Petroleum, biological studies
     Polyamides, biological studies
     Polyesters, biological studies
     Polyoxyalkylenes, biological studies
Polysaccharides, biological studies
     Polysiloxanes, biological studies
     Proteins
     Salts, biological studies
     Shellac
     Soybean oil
     Tannins
     Waxes
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (targeted controlled delivery microspheres activated by changes in pH
        or salt concentration)
·IT
     Amines, biological studies
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (tertiary; targeted controlled delivery microspheres activated by
        changes in pH or salt concentration)
     Fats and Glyceridic oils, biological studies
IT
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (vegetable, hydrogenated; targeted controlled delivery microspheres
        activated by changes in pH or salt concentration)
     Lactams
     RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
         (β-, antibiotics; targeted controlled delivery microspheres
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activated by changes in pH or salt concentration) IT Antibiotics (β-lactam; targeted controlled delivery microspheres activated by changes in pH or salt concentration) 18479-58-8, Dihydromyrcenol 78-70-6, Linalool 106-24-1, Geraniol IT 20780-48-7, Tetrahydrolinalyl acetate RL: COS (Cosmetic use); MSC (Miscellaneous); BIOL (Biological study); USES (targeted controlled delivery microspheres activated by changes in pH or salt concentration) 50-70-4, Sorbitol, biological studies 56-81-5, Glycerin, biological IT 57-48-7, Fructose, biological studies 57-55-6, Propylene glycol, biological studies 69-72-7D, Salicylic acid, benzophenyl derivs. 69-72-7D, Salicylic acid, derivs. 69-93-2, Uric acid, biological studies 79-10-7D, Acrylic acid, derivs., polymers 79-41-4D, Methacrylic acid, derivs., polymers 87-99-0, Xylitol 91-64-5D, Coumarin, derivs. 94-41-7, Benzalacetophenone 107-21-1, Ethylene glycol, biological 107-41-5, Hexylene glycol 107-64-2, Distearyl dimethyl ammonium chloride 107-88-0, 1,3-Butanediol 118-55-8, Salol 118-92-3D, Anthranilic acid, derivs. 119-61-9D, Benzophenone, ammonium chloride 119-61-9D, Benzophenone, derivs. 123-31-9, Hydroquinone, biological studies 150-13-0, p-Aminobenzoic acid 538-58-9, Dibenzalacetone 621-82-9D, Cinnamic acid, derivs. 2568-33-4. Isoprene glycol 3843-16-1, Distearyldimethylammonium methylsulfate 9002-88-4, Polyethylene 9002-89-5, Polyvinyl alcohol 9003-01-4, Polyacrylic acid 9004-34-6, Cellulose, biological studies Cellulose, oxidized 9004-35-7, Cellulose acetate 9005-25-9004-34-6D. 9005-25-8D, Starch, 9006-26-2, Ethylene-maleic anhydride copolymer 9006-65-9, derivs. 9011-16-9, Methyl vinyl ether-maleic anhydride copolymer Dimethicone, 17301-53-0, Behenyltrimethylammonium chloride 24937-49-3, Polyornithine 24938-16-7, Eudragit EPO 25085-34-1, Acrylic acid-styrene 25104-12-5, Polyornithine 25104-18-1, Polylysine copolymer 25300-64-5, Maleic acid-styrene copolymer 25322-68-3, Polyethylene oxide 25322-69-4, Polypropylene glycol 25609-89-6, Crotonic acid-vinyl acetate copolymer 26336-38-9, Polyvinylamine 27323-69-9D, Dihydroxycinnamic 38000-06-5, Polylysine 52907-01-4, Cellulose acetate acid, derivs. trimellitate 65497-29-2, Guar hydroxypropyl trimonium chloride 66829-29-6, Hi-Cap 100 71138-97-1, Hydroxypropyl methyl 220828-91-1, Incroquat Behenyl HE cellulose acetate succinate 426258-09-5, Incrosoft 100 RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (targeted controlled delivery microspheres activated by changes in pH or salt concentration) 50-02-2, Dexamethasone 50-23-7, Hydrocortisone 50-24-8, Prednisolone 50-33-9, Phenylbutazone, biological studies 50-78-2, Aspirin Cortisone 53-86-1, Indomethacin 54-64-8, Thimerosal 55-56-1, Chlorhexidine 55-86-7, Chloramine 56-75-7, Chloramphenicol 60-54-8, Tetracycline 60-80-0, Antipyrine 75-47-8, Iodoform Procaine 94-09-7, Benzocaine 108-95-2, Phenol, biological studies 114-07-8, Erythromycin 124-94-7, Triamcinolone 137-58-6, Lidocaine Fluocortolone 378-44-9, β-Methasone 443-48-1, Metronidazole 152-97-6, 1404-04-2, Neomycin 1404-26-8, Polymyxin B 1404-90-6, Vancomycin 1405-97-6, Gramicidin 1406-05-9, Penicillin 5104-49-4, Flurbiprofen 8025-81-8, Spiramycin 8063-07-8, Kanamycin 10043-35-3, Boric acid, biological studies 15687-27-1, Ibuprofen 20283-69-6 22204-53-1, 32385-11-8, Sisomicin 32986-56-4, Tobramycin Naproxen **35607-66-0**, Cefoxitin 36322-90-4, Piroxicam 37517-28-5, 64221-86-9 Amikacin 59995-64-1, Thienamycin RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (targeted controlled delivery microspheres activated by changes in pH or salt concentration) 65497-29-2, Guar hydroxypropyl trimonium chloride RL: COS (Cosmetic use); MSC (Miscellaneous); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (targeted controlled delivery microspheres activated by changes in pH or salt concentration) **35607-66-0**, Cefoxitin IT

RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)

or salt concentration)

(targeted controlled delivery microspheres activated by changes in pH

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ANSWER 2 OF 6 HCAPLUS COPYRIGHT 2005 ACS on STN
AN
     2003:42092 HCAPLUS
DN
     138:112443
ED
     Entered STN: 17 Jan 2003
     Tablet compositions for poorly-compressible pharmaceuticals
ΤI
     Matharu, Amol Singh; Patel, Mahendra R.
IN
     Geneva Pharmaceuticals, Inc., USA
PA
     PCT Int. Appl., 20 pp.
S<sub>0</sub>
     CODEN: PIXXD2
DT
     Patent
     English
LA
     ICM A61K031-155
     ICS
         A61K009-20
CC
     63-6 (Pharmaceuticals)
FAN. CNT 1
     PATENT NO.
                                                                      DATE
                          KIND
                                 DATE
                                              APPLICATION NO.
                                                                      20020627 <---
PΙ
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                                 20030116
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     US 2003021841
                           A1
                                              US 2002-183881
PRAI US 2001-302613P
                                 20010702
CLASS
                         PATENT FAMILY CLASSIFICATION CODES
 PATENT NO.
                  CLASS
 WO 2003004009
                  ICM
                         A61K031-155
                         A61K009-20
                  ICS
 US 2003021841
                  ECLA
                         A61K009/20H4; A61K009/20H6F2
     The present invention relates to a process for preparing tablet dosage forms
     of poorly-compressible pharmaceuticals and to tablet dosage forms. The
     process is especially useful for preparing tablets of the poorly-compressible drug
     metformin-HCl. Thus, tablets contained metformin-HCl 500, HPMC 320, stearyl alc. 200, and Mg stearate mg/unit.
     tablet poorly compressible pharmaceutical
     Glycerides, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (C16-18, hydrophobic excipient; tablet compns. for poorly-compressible
        pharmaceuticals)
IT
     Fatty acids, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (esters, hydrophobic excipient; tablet compns. for poorly-compressible
        pharmaceuticals)
IT
     Alcohols, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (fatty, hydrophobic excipient; tablet compns. for poorly-compressible
         pharmaceuticals)
     Polyoxyalkylenes, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (hydrophilic excipient; tablet compns. for poorly-compressible
         pharmaceuticals)
IT
     Fatty acids, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (hydrophobic excipient; tablet compns. for poorly-compressible
         pharmaceuticals)
     Protamines
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (sulfates; tablet compns. for poorly-compressible pharmaceuticals)
     Dissolution
     Hypnotics and Sedatives
         (tablet compns. for poorly-compressible pharmaceuticals)
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IT Carnauba wax RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (tablet compns. for poorly-compressible pharmaceuticals) IT Drug delivery systems (tablets; tablet compns. for poorly-compressible pharmaceuticals) 63-42-3, Lactose 9000-01-5, Acacia gum **9000-30-0, Guar** gum 9000-65-1, Tragacanth gum 9003-01-4, Carboxypolymethylene ΙT 9003-39-8, Polyvinylpyrrolidone 9004-32-4, Carboxymethyl cellulose sodium salt 9004-34-6D, Cellulose, derivs. 9004-42-6, Carboxyethyl cellulose 9004-62-0, Hydroxyethyl cellulose 9004-64-2, Hydroxyethyl cellulose 9004-62-0, Hydroxyethyl cellulose 9004-65-3, HPMC 9004-67-5, Methyl 9004-67-5, Methyl cellulose 9005-38-3, cellulose Sodium alginate 11138-66-2, Xanthan gum 37353-59-6, Hydroxymethyl cellulose 25322-68-3, Polyethylene glycol 74811-65-7, Croscarmellose sodium RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hydrophilic excipient; tablet compns. for poorly-compressible pharmaceuticals) 57-11-4, Stearic acid, biological studies 79-41-4D, Methacrylic acid, IT 112-85-6, Behenic acid 112-92-5, Stearyl alcohol 9004-57-3, polymers Ethyl cellulose 9010-88-2, Eudragit NE 30D 31566-31-1, Glyceryl monostearate 36653-82-4, Cetyl alcohol 77538-19-3, Glyceryl behenate RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (hydrophobic excipient; tablet compns. for poorly-compressible pharmaceuticals) ΙT 9004-34-6, Cellulose, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (microcryst.; tablet compns. for poorly-compressible pharmaceuticals) 50-02-2, Dexamethasone 50-24-8, Prednisolone 50-33-9, Phenylbutazone, ΙT biological studies 50-34-0, Propantheline bromide 50-47-5, Desmethylimipramine 50-48-6, Amitriptyline 50-49-7, Imipramine 50-52-2, Thioridazine 50-53-3, Chlorpromazine, biological studies 50-54-4, Quinidine sulfate 50-78-2, Acetylsalicylic acid 50-81-7, Ascorbic acid, biological studies 51-06-9, Procainamide 51-34-3, Scopolamine 51-43-4, Epinephrine 51-48-9, Thyroxine, biological studies 51-52-5, Propylthiouracil 52-01-7, Spironolactone 52-28-8, Calaina absorbate 52-52-6, Varappril 52-68-8, Haloparidel 53-03-2 Codeine phosphate 52-53-9, Verapamil 52-86-8, Haloperidol 53-03-2, Prednisone 53-86-1, Indomethacin 54-31-9, Furosemide 55-56-1, Chlorhexidine 55-63-0, Glyceryl trinitrate 57-27-2, Morphine, biological studies 57-42-1, Pethidine 57-66-9, Probenecid 57-96-5, Sulfinpyrazone 58-55-9, Theophylline, biological studies 58-73-1, Diphenhydramine 58-74-2, Papaverine 58-93-5, Hydrochlorthiazide 59-02-9 59-30-3, Folic acid, biological studies 59-43-8, Thiamine, biological studies 59-67-6, Nicotinic acid, biological studies 60-13-9, Amphetamine sulfate 60-32-2, &-Aminocaproic acid 61-68-7, Mefenamic acid 61-75-6, Bretylium tosylate 64-75-5, Tetracycline hydrochloride 64-77-7, Tolbutamide 65-23-6, Pyridoxine 69-23-8, Fluphenazine 69-53-4, Ampicillin 76-42-6, Oxycodone 77-19-0, Dicyclomine 77-36-1, Chlorthalidone 78-11-5, Pentaerythritol tetranitrate 86-22-6, Brompheniramine 86-54-4, Hydralazine 87-33-2, Isosorbide dinitrate 90-82-4, Pseudoephedrine 93-14-1, Guaiphenesin 94-20-2, Diabinese 103-90-2, Acetaminophen 113-15-5, Ergotamine 113-45-1, Methylphenidate 113-92-8 117-89-5, Trifluperazine 122-09-8, Phentermine 123-03-5, Cetylpyridinium chloride 125-71-3, Dextromethorphan 128-62-1, Noscapine 129-06-6, Warfarin sodium 134-80-5, Diethylpropion hydrochloride 146-22-5, Nitrazepam 147-20-6, Diphenylpyraline 299-28-5, Calcium gluconate 299-42-3, Ephedrine 315-30-0, Allopurinol 317-34-0, Aminophylline 318-98-9, Propranolol hydrochloride 364-62-5, Metoclopramide 389-08-2, Nalidixic acid 396-01-0, Triamterene 404-82-0, Fenfluramine hydrochloride 439-14-5, Diazepam 456-59-7, Cyclandelate 480-30-8, Dichloralphenazone 554-13-2, Lithium carbonate 555-30-6, Methyldopa 561-27-3, Heroin 577-11-7, Dioctylsodium sulfosuccinate 587-23-5, Methenamine mandelate 577-11-7, Dioctylsodium sulfosuccinate 587-23-5, Methenamine mandelate 603-50-9, Bisacodyl 630-93-3, Phenytoin sodium 638-23-3, Carbocisteine 645-43-2, Guanethidine monosulfate 643-22-1, Erythromycin stearate 846-50-4, Temazepam 915-30-0, Diphenoxylate 1069-66-5, Sodium valproate 1115-70-4, Metformin hydrochloride 1309-42-8, Magnesium hydroxide 1335-30-4, Aluminum silicate 1404-88-2, Tyrothricin 1420-53-7, Codeine sulfate 1617-90-9, Vincamine 1668-19-5, Doxepin 1847-24-1, Flucloxacillin sodium 2706-50-5, Amphetamine hydrochloride

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3200-06-4, Naftidrofuryl oxalate 3572-80-3, Cyclazocine
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                                                            7054-25-3, Quinidine
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42399-41-7, Diltiazem
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      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (tablet compns. for poorly-compressible pharmaceuticals)
                THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE. CNT
RE
(1) Abdallah, 0; S T P PHARMA 1988, V4(1), P15 HCAPLUS
    Depomed Inc; WO 9855107 A 1998 HCAPLUS
    Kumar, T; WO 0228181 A 2002 HCAPLUS
Kumar, V; US 6117451 A 2000 HCAPLUS
    Squibb Bristol Myers Co; WO 9947128 A 1999 HCAPLUS
     9000-30-0, Guar gum 11138-66-2,
     Xanthan gum
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (hydrophilic excipient; tablet compns. for poorly-compressible
         pharmaceuticals)
ΙT
      9010-88-2, Eudragit NE 30D
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (hydrophobic excipient; tablet compns. for poorly-compressible
         pharmaceuticals)
      15686-71-2, Cephalexin
IT
      RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (tablet compns. for poorly-compressible pharmaceuticals)
L81 ANSWER 3 OF 6 HCAPLUS COPYRIGHT 2005 ACS on STN
AN
      2002:555334 HCAPLUS
DN
      137:114525
ED
      Entered STN: 26 Jul 2002
      Syntactic deformable pharmaceutical foam compositions
ΤI
      Odidi, Isa; Odidi, Amina
IN
PA
     Can.
S<sub>0</sub>
     PCT Int. Appl., 47 pp.
      CODEN: PIXXD2
DT
     Patent
     English
LA
      ICM A61K009-00
IC
CC
     63-6 (Pharmaceuticals)
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PRAI US 2001-765783
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     WO 2002-CA54
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CLASS
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 WO 2002056861
                         A61K009-00
                  T CM
                         A61K009/20K2B; A61K009/20P
US 6800668
                  ECLA
    The invention relates to methods for preparing a syntactic foam composition
     suitable for use as a carrier for chems. or other compds., including
     pharmaceuticals. Carbopol 971P, hydroxyethyl cellulose, cellulose
     microspheres and silica, was mixed in a high-shear mixer. The resulting admixt. was treated with 2-propanol, while simultaneously subjecting the
     admixt, to high-shear forces in the high-shear mixer. This mixing created
     a uniform stable syntactic deformable and compressible dendritic solid foam which could be shaped before drying. Metoprolol succinate was added
     to the above admixt. and subjected to high-shear agitation for 2 min
     before treatment with 2-propanol. A stable syntactic deformable and
     compressible dendritic solid foam which could be shaped before drying was
     obtained. This was dried at 40°. The dried foam was the
     disentangled by size reduction to obtain discrete particles.
                                                                       The free flowing
     particles were reassembled and shaped by compression in a mold. The
     shaped units, when subjected to an aqueous medium, released metoprolol over a
     period of ≤3 h.
     syntactic deformable foam drug delivery; polymer syntactic deformable foam
     drug delivery
     Polyesters, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (caprolactone-based; syntactic deformable pharmaceutical foam compns.)
     Vinyl compounds, biological studies
IT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (carboxy-containing, polymers; syntactic deformable pharmaceutical foam
        compns.)
     Drug delivery systems
IT
         (controlled-release; syntactic deformable pharmaceutical foam compns.)
     Drug delivery systems
IT
         (foams; syntactic deformable pharmaceutical foam compns.)
IT
     Polyesters, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (hydroxycarboxylic acid-based; syntactic deformable pharmaceutical foam
         compns.)
     Polyesters, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (lactic acid-based; syntactic deformable pharmaceutical foam compns.)
     Drug delivery systems
IT
         (microspheres; syntactic deformable pharmaceutical foam compns.)
ΙT
     Coating materials
     Human
     Size reduction
     Stabilizing agents
         (syntactic deformable pharmaceutical foam compns.)
     Metals, biological studies
     Polyesters, biological studies
     Polyoxyalkylenes, biological studies
     Polysaccharides, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
         (syntactic deformable pharmaceutical foam compns.)
     39391-18-9, Cyclooxygenase
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     124-38-9, Carbon dioxide, processes 7727-37-9, Nitrogen, processes
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     process); PROC (Process)
         (liquid; syntactic deformable pharmaceutical foam compns.)
     329900-75-6, Cyclooxygenase-2
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
         (syntactic deformable pharmaceutical foam compns.)
     67-63-0, 2-Propanol, uses
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)
         (syntactic deformable pharmaceutical foam compns.)
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50-02-2, Dexamethasone 50-28-2, Estradiol, biological studies 50-48-6, Amitriptyline 50-70-4, Sorbitol, biological studies 50-78-2, Aspirin 50-99-7, Glucose, biological studies 51-48-9, Levothyroxine, biological studies 53-03-2, Prednisone 54-31-9, Furosemide 57-27-2, Morphine, 57-41-0, Phenytoin 57-50-1, Sucrose, biological biological studies studies 57-63-6, EthinylEstradiol 58-93-5, Hydrochlorothiazide 59-92-7, Levodopa, biological studies 60-87-7, Promethazine 63-42-3, Lactose 67-20-9, Nitrofurantoin 68-22-4, Norethindrone 69-65-8, Mannitol 76-42-6, Oxycodone 76-57-3, Codeine 78-44-4, Carisoprodol 81-81-2, Warfarin 83-43-2, Methylprednisolone 87-99-0, Xylitol 89-57-6, Mesalamine 90-82-4, Pseudoephedrine 93-14-1, Guaifenesin 99-66-1, Pentangic acid 2-propyl 103-00-2 Acataminoshop 114-07-9 99-66-1, Pentanoic acid, 2-propyl 103-90-2, Acetaminophen 114-07-8, Erythromycin 125-29-1, Hydrocodone 127-07-1, Hydroxyurea 132-98-9, Popisillip VK 155-00-0 Translamentary 200-69-00 American Erythromycin 125-29-1, Hydrocodone 127-07-1, Hydroxyurea 132-98-9, Penicillin VK 155-09-9, Tranylcypromine 300-62-9D, Amphetamine, salts 303-53-7, Cyclobenzaprine 315-30-0, Allopurinol 378-44-9, Betamethasone 396-01-0, Triamterene 439-14-5, Diazepam 469-62-5, Propoxyphene 525-66-6, Propranolol 673-06-3, D-Phenylalanine 797-63-7, Levonorgestrel 846-49-1, Lorazepam 846-50-4, Temazepam 1119-34-2, L-Arginine hydrochloride 1622-61-3, Clonazepam 3056-17-5, Stavudine 3930-20-9 Sotalol 4205-20-7 Clonidina 4410-20-0 Stavudine 3930-20-9, Sotalol 4205-90-7, Clonidine 4419-39-0, Beclomethasone 7447-40-7, Potassium Chloride, biological studies 7460-12-0, Pseudoephedrine sulfate 7481-89-2, Zalcitabine 7631-86-9, Silica, biological studies 9002-89-5, Polyvinyl alcohol 9002-96-4, α-Tocopherol polyethylene glycol succinate 9003-39-8, Povidone 9004-34-6, Cellulose, biological studies 9004-54-0, Dextran, biological studies 9004-62-0, Hydroxyethyl Cellulose 9004-65-3, Hydroxypropyl Methyl cellulose 9005-25-8, Starch, biological studies Calcitonin 10238-21-8, Glyburide 10540-29-1, Tamoxifen Xanthan gum 12650-69-0, 11138-66-2 Mupirocin **15686-71-2**, Cephalexin 15687-27-1, Ibuprofen 16051-77-7, Isosorbide Mononitrate 18559-94-9, Albuterol Glyceryl behenate 19794-93-5, Trazodone 20830-75-5, Digoxin 21256-18-8, Oxaprozin 22204-53-1, Naproxen 23593-75-1, Clotrimazole 24980-41-4, Poly(e-caprolactone) 25086-15-1, Eudragit 25248-42-4, Poly[oxy(1-oxo-1, 6-hexanediyl)] 25322-68-3, Polyethylene glycol 25812-30-0, Gemfibrozil 26009-03-0, Poly(glycolic acid) 26023-30-3, Poly[oxy(1-methyl-2-oxo-1, 2-ethanediyl)] 26100-51-6, Poly(lactic acid) 26124-68-5, Poly(glycolic acid) 26787-78-0, 26100-51-6, Amoxicillin 28860-95-9, Carbidopa 28981-97-7, Alprazolam 29122-68-Atenolol 30516-87-1, Zidovudine 32986-56-4, Tobramycin 34346-01-5, Glycolic acid-lactic acid copolymer 51384-51-1, Metoprolol 54739-18-29122-68-7, 54739-18-3, Fluvoxamine 54910-89-3, Fluoxetine 55268-75-2, Cefuroxime 56180-94-0, Acarbose 58001-44-8 59122-46-2, Misoprostol 59729-33-8. Citalopram 59803-98-4, 61869-08-7, Paroxetine 59803-98-4, Brimonidine 60205-81-4, Ipratropium 63590-64-7, Terazosin 63675-72-9, Nisoldipine 66376-36-1, Alendronate 66722-44-9, Bisoprolol 66357-35-5, Ranitidine 00507-35-5, Kanitidine 00570-36-1, Alendronate 60722-44-9, Bisoprolol 69655-05-6, Didanosine 72432-03-2, Miglitol 72509-76-3, Felodipine 72956-09-3, Carvedilol 74191-85-8, Doxazosin 75330-75-5, Lovastatin 75847-73-3, Enalapril 76547-98-3, Lisinopril 76584-70-8, Divalproex sodium 76824-35-6, Famotidine 76963-41-2, Nizatidine 78644-42-5, Poly (malic acid) 78666-19-0, Poly (malic acid), SRU 79617-96-2, Sertraline 79794-75-5, Loratadine 79902-63-9, Simvastatin 81098-60-4, Cicappide 81102-11-9 Clasitherania 81093-37-0, Pravastatin 81098-60-4, Cisapride 81103-11-9, Clarithromycin 82419-36-1, Ofloxacin 82626-48-0, Zolpidem 83799-24-0, Fexofenadine 83881-51-0, Cetirizine 83905-01-5, Azithromycin 84449-90-1, Raloxifene 85441-61-8, Quinapril 85721-33-1, Ciprofloxacin 86541-75-5, Benazepril 87333-19-5, Ramipril 82626-48-85721-33-1, Ciprofloxacin 89365-50-4, Salmeterol 91161-71-6, Terbinafine 93413-69-5, Venlafaxine 93957-54-1, Fluvastatin 97322-87-7, 88150-42-9, Amlodipine 92665-29-7, Cefprozil 93479-97-1, Glimepiride 98048-97-6, Fosinopril 98418-47-4, Metoprolol succinate dansetron 100986-85-4, Levofloxacin 103577-45-3, Troglitazone 99614-02-5, Ondansetron 103628-46-2, Sumatriptan 104632-26-0, Pramipexole Lansoprazole 106133-20-4, Tamsulosin 106266-06-2, 105102-22-5, Mometasone 109889-09-0, Granisetron 107753-78-6, Zafirlukast Risperidone 113665-84-2, Clopidogrel 124937-51-5, Tolterodine 111974-69-7, Quetiapine 114798-26-4, Losartan 127779-20-8. 120014-06-4, Donepezil Saquinavir 129618-40-2, Nevirapine 130209-82-4, Latanoprost 134523-00-5, Atorvastatin 134678-17-4, Repaglinide 136470-78-5, Abacavir 132539-06-1, Olanzapine 134523-00-5, Lamivudine 135062-02-1, Repaglinide

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136817-59-9, Delayirdine 137862-53-4, Valsartan
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        (syntactic deformable pharmaceutical foam compns.)
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L81
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AN
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     Entered STN: 10 Aug 2001
ΤI
     Shell-and-core dosage form approaching zero-order drug release
     Berner, Bret; Louie-Helm, Jenny; Gusler, Gloria; Shell, John N.
IN
PA
     Depomed, Inc., USA
S0
     PCT Int. Appl., 36 pp.
     CODEN: PIXXD2
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     English
     ICM A61K009-00
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     63-6 (Pharmaceuticals)
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 US 2003104062
                  ECLA
                          A61K009/20H6B; A61K009/28H6B
     Drugs are formulated as oral dosage forms for controlled release in which
     the release rate limiting portion is a shell surrounding the drug-containing
     core. The shell releases drug from the core by permitting diffusion of
     the drug from the core. The shell also promotes gastric retention of the
     dosage form by swelling upon imbibition of gastric fluid to a size that is
     retained in the stomach during the postprandial or fed mode. Thus, core
     containing Polyox-303 700 and the shell 200 mg was prepared with the drug
     loading in the core being 71.4% by weight (with no drug contained in the
     shell). The release rate approached zero order.
     zero order controlled drug release; polymer core controlled drug release
     Drug delivery systems
         (capsules, controlled-release; shell-and-core dosage form approaching
        zero-order drug release)
ΙT
     Drug delivery systems
         (controlled-release; shell-and-core dosage form approaching zero-order
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drug release)
IT
     Diffusion
     Dissolution rate
     Molecular weight distribution
        (shell-and-core dosage form approaching zero-order drug release)
     Polymers, biological studies
     Polyoxyalkylenes, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (shell-and-core dosage form approaching zero-order drug release)
IT
     Drug delivery systems
        (tablets, controlled-release; shell-and-core dosage form approaching
        zero-order drug release)
IT
     9003-01-4, Polyacrylic acid
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (crosslinked; shell-and-core dosage form approaching zero-order drug
        release)
IT
     1115-70-4, Metformin hydrochloride
     RL: BPR (Biological process); BSU (Biological study, unclassified); THU
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        (shell-and-core dosage form approaching zero-order drug release)
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     3847-29-8, Erythromycin lactobionate 7439-89-6D, Iron, salts, biological
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         (shell-and-core dosage form approaching zero-order drug release)
     11138-66-2, Xanthan gum 64544-07-6,
     Cefuroxime axetil
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     2000:900423 HCAPLUS
AN
DN
     134:61525
ED
     Entered STN: 22 Dec 2000
     Taste masked compositions comprising methacrylic acid copolymer and
ΤI
     phthalate polymer
     Mukherji, Gour; Goel, Sandhya; Arora, Vinod Kumar
     Ranbaxy Laboratories Limited, India
PA
S0
     PCT Int. Appl., 19 pp.
     CODEN: PIXXD2
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     English
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     A61K009-16; A61K009-22; A61K047-30
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     63-6 (Pharmaceuticals)
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                                                                A61K047-30
 US 6565877
                          A61K009/16H6F; A61K009/16H6B
                  ECLA
     A taste masked composition which comprises a bitter tasting drug, a combination
     of two enteric polymers comprising, a methacrylic acid copolymer and a
     phthalate polymer is described. The composition of the present invention is
     prepared by dissolving the active ingredient, the methacrylic acid copolymer
     and the phthalate polymer in a solvent and recovering the composition from the
     solution thereof. Cefuroxime axetil 2 g were blended with 2 g
     Eudragit L100-55/hydroxypropyl Me cellulose phthalate (7:3) and
     dissolved in 20 mL acetone containing 5 % water.
                                                            The resulting mixture was
     dried and sized to obtain taste-masked granules.
     taste masked oral polymethacrylate cellulose phthalate; oral granule
     cefuroxime Eudragit cellulose phthalate
IT
     Drug delivery systems
         (granules, oral; taste masked compns. comprising methacrylic acid
         copolymer and phthalate polymer).
IT
     Antibiotics
         (macrolide; taste masked compns. comprising methacrylic acid copolymer
         and phthalate polymer)
ΙT
     Antibiotics
         (quinolone; taste masked compns. comprising methacrylic acid copolymer
         and phthalate polymer)
IT
     Drug delivery systems
         (suspensions; taste masked compns. comprising methacrylic acid
         copolymer and phthalate polymer)
IT
     Drug delivery systems
         (syrups; taste masked compns. comprising methacrylic acid copolymer and
         phthalate polymer)
IT
     Drug delivery systems
         (tablets; taste masked compns. comprising methacrylic acid copolymer
         and phthalate polymer)
     50-53-3, Chlorpromazine, biological studies 56-75-7, Chloramphenicol 63-42-3, Lactose 69-65-8, D-Mannitol 80-62-6D, Methyl methacrylate, copolymers 114-07-8, Erythromycin 7647-14-5, Sodium chloride,
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     biological studies
     acetate phthalate
     9063-38-1, Sodium starch glycolate 11111-12-9, Cephalosporin
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     11138-66-2, Xanthan gum
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         (taste masked compns. comprising methacrylic acid copolymer and
         phthalate polymer)
7 THERE ADD 7
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    11138-66-2, Xanthan gum 55268-75-2,
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     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (taste masked compns. comprising methacrylic acid copolymer and
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L81 ANSWER 6 OF 6 HCAPLUS COPYRIGHT 2005 ACS on STN
     1998:268331 HCAPLUS
AN
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     Entered STN: 11 May 1998
ED
     Pharmaceutical composition for rapid suspension in aqueous media
TI
     Calanchi, Massimo Maria; Marconi, Marco Giuseppe Raffaele; Mapelli, Luigi
IN
     Giovanni
PA
     Eurand International S.P.A., Italy
S<sub>0</sub>
     PCT Int. Appl., 30 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
     ICM A61K009-00
IC
         A61K009-20; A61K009-16
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CLASS
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                  ICM
                         A61K009-20; A61K009-16
A61K009/00N2
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 GB 2318511
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                  ECLA
 US 6261602
     The invention provides a granular composition useful as a pharmaceutical
     carrier which can be used for the preparation of pharmaceutical compns. that
     are capable of rapid suspension in water or aqueous media including saliva.
     The compns. may be used by addition to a glass of water with stirring or
     taken directly in the mouth. The granular composition may be prepared by a
     process which comprises subjecting a mixture of a thickening agent and a
     disintegrating agent to wet granulation with an aqueous medium as wetting
     agent or dry granulation to make a novel granular product and preparing the
     pharmaceutical composition from the granular product and the drug. A
     water-soluble inert excipient, which may be a sugar, may be mixed with the
     granular product prior to mixing with the drug. Base granules were prepared containing Keltrol F, Ac-di-Sol, Avicel PH 200 and Explotab. These
     granules were mixed with Karion, aspartame and orange flavor and monodose
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sachets were prepared from this mixture and 5-aminosalicylic acid coated with
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         pharmaceutical granule suspension
         Drug delivery systems
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ΙT
         Buffers
         Lubricants
         Sweetening agents
         Thickening agents
         Vitamins
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                               9000-36-6, Karaya gum
         Guar gum
         9000-65-1, Gum tragacanth 9002-18-0, Agar
                                                                                                    9005-25-8, Starch,
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                                                                                                     9005-38-3, Sodium alginate
         biological studies
         9050-04-8, Calcium carboxymethyl cellulose 9050-36-6, 9063-38-1, Sodium starch glycolate 11138-66-2, Xanthan
                                                                                                9050-36-6, Maltodextrin
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         90-82-4, Pseudoephedrine 93-14-1, Guaifenesin 103-90-2, Paracetamol 114-07-8, Erythromycin 125-71-3, Dextromethorphan 364-62-5,
         Metoclopramide 439-14-5, Diazepam 554-13-2, Lithium carbonate 616-91-1, Acetylcysteine 638-23-3 1406-05-9, Penicillin 1812-30-2,
                                  3820-67-5, Glafenine 5250-39-5, Flucloxacillin 8049-47-6, 11111-12-9, Cephalosporin 14838-15-4, Phenylpropanolamine
         Bromazepam
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         15307-86-5, Diclofenac 15686-71-2, Cefalexin 15687-27-1,
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25812-30-0, Gemfibrozil
41340-25-4, Etodolac
         Ibuprofen 16051-77-7, Isosorbide mononitrate 19216-56-9, Prazosin 22071-15-4, Ketoprofen
         26787-78-0, Amoxicillin 31637-97-5, Etotibrate 41040 20 4, 20040142 42399-41-7, Diltiazem 51481-61-9, Cimetidine 54910-89-3, Fluoxetine 66357-35-5, Ranitidine
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         RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
                (pharmaceutical composition for rapid suspension in aqueous media)
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         9000-30-0, Guar gum 11138-66-2,
         Xanthan gum
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         15686-71-2, Cefalexin
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